



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>6</sup> :</b> <b>C12Q</b>	<b>A2</b>	<b>(11) International Publication Number:</b> <b>WO 97/18323</b> <b>(43) International Publication Date:</b> 22 May 1997 (22.05.97)
<b>(21) International Application Number:</b> PCT/US96/19337 <b>(22) International Filing Date:</b> 18 November 1996 (18.11.96)  <b>(30) Priority Data:</b> 08/558,666 16 November 1995 (16.11.95) US 08/007,312 27 February 1996 (27.02.96) US 08/725,304 21 October 1996 (21.10.96) US  <b>(71) Applicant (for all designated States except US):</b> ICOS CORPORATION [US/US]; 22021 20th Avenue, S.E., Bothell, WA 98021 (US).  <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> HOEKSTRA, Merl, F. [US/US]; 10321 216th Street, S.E., Snohomish, WA 98290 (US). HOLTZMAN, Doug, A. [US/US]; 4308 4th Avenue, N.E., Seattle, WA 98105 (US). KEEGAN, Kathleen, S. [US/US]; 5812 Mercer Way, Mercer Island, WA 98040 (US).  <b>(74) Agent:</b> NOLAND, Greta, E.; Marshall, O'Toole, Gerstein, Murray & Borun, 6300 Sears Tower, 233 South Wacker Drive, Chicago, IL 60606-6402 (US).		<b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>Without international search report and to be republished upon receipt of that report.</i>
<b>(54) Title:</b> CELL CYCLE CHECKPOINT PIK-RELATED KINASE MATERIALS AND METHODS		
<b>(57) Abstract</b> <p>The present invention generally relates to genes encoding cell cycle checkpoint phosphatidylinositol kinase (PIK)-related proteins essential to DNA damage responses in cells. These PIK-related kinases are required in regulatory pathways that arrest the cell cycle following DNA damage to allow DNA repair prior to mitosis or initiation of DNA replication. More particularly, the invention provides a novel human cell cycle checkpoint PIK-related kinase, MCCS1, and polynucleotide sequences encoding the MCCS1. Assays for identifying modulators of MCCS1 useful as, for example, chemotherapy and radiation adjuvants, are also provided by the invention. Further, assays for identifying modulators of the cell cycle checkpoint phosphatidylinositol kinase (PIK)-related protein identified as ATM are provided.</p>		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

## CELL CYCLE CHECKPOINT PIK-RELATED KINASE MATERIALS AND METHODS

### FIELD OF THE INVENTION

5 The present invention generally relates to genes encoding cell-cycle  
checkpoint phosphatidylinositol kinase (PIK)-related genes and proteins essential to  
DNA damage responses in cells. The checkpoint kinases play a role in the  
surveillance of DNA damage that occurs as a result of replication errors, DNA  
mismatches, radiation treatment, or chemotherapeutic drugs. These kinases are  
required in regulatory pathways that lead to cell cycle arrest following DNA damage,  
10 giving the cell notice and time to correct lesions prior to the initiation of DNA  
replication. More particularly, the invention relates to a novel human PIK-related  
kinase, Mammalian Cell Cycle Surveillance 1 (MCCS1), polynucleotides encoding  
the PIK-related kinase, and methods for assaying and modulating the enzymatic  
activity of the kinase and related kinases.

### BACKGROUND

15 The process of eukaryotic cell growth and division is the somatic or  
mitotic cell cycle which consists of four phases, the G<sub>1</sub> phase, the S phase, the G<sub>2</sub>  
phase, and the M phase. The G<sub>1</sub>, S, and G<sub>2</sub> phases are collectively referred to as  
interphase of the cell cycle. The cell cycle is structurally and functionally conserved  
20 in its basic process and mode of regulation across all eukaryotic species. During the  
G<sub>1</sub> (gap) phase, biosynthetic activities of the cell progress at a high rate. The S  
(synthesis) phase begins when DNA synthesis starts and ends when the DNA content  
of the nucleus of the cell has been replicated and two identical sets of chromosomes  
are formed. The cell then enters the G<sub>2</sub> (gap) phase which continues until mitosis  
25 starts. In mitosis, the chromosomes pair and separate and two new nuclei form, and  
in cytokinesis the cell itself splits into two daughter cells each receiving one nucleus

-2-

containing one of the two sets of chromosomes. Mitosis and cytokinesis together form the M (mitosis) phase of the cell cycle. Cytokinesis terminates the M phase and marks the beginning of interphase of the next cell cycle. The sequence in which the events in the cell cycle proceed is tightly regulated such that the initiation of one cell cycle event is dependent on the completion of the prior cell cycle event. This allows fidelity in the duplication and segregation of genetic material from one generation of cells to the next.

The term "cell cycle checkpoints" refers to the proteins, signals, processes, and feedback controls that integrate discontinuous events during cellular replication, in order to maintain essential dependencies within the cell cycle. The present invention specifically relates to the cell cycle checkpoint that ensures that mitosis is delayed until the completion of DNA synthesis and/or the accurate repair of DNA damage occurs.

Failure of cell cycle checkpoints predisposes individuals to or directly causes many disease states such as cancer, ataxia telangiectasia, embryo abnormalities, and various immunological defects associated with aberrant B and T cell development. The latter are associated with pathological states such as lupus, arthritis and autoimmune diseases. Intense research efforts have therefore focused on identifying cell cycle checkpoints and the proteins essential for the function of the checkpoints.

Genetic analysis in the yeasts *Schizosaccharomyces pombe* and *Saccharomyces cerevisiae* has identified a number of genes important for cell cycle arrest and DNA repair responses to ionizing radiation (IR). For a review, see Carr and Hoekstra, *Trends in Cell Biology*, 5: 32-40 (1995). One such gene, identified in both yeasts, is required for a DNA damage checkpoint which arrests the cell cycle at the G2 phase, as well as a related checkpoint which monitors the completion of DNA synthesis and arrests the cell cycle at the S phase. The gene is named *rad3+* in *S. pombe* [Seaton *et al.*, *Gene*, 119: 83-89 (1992)], *MEC1/ESR1* in *S. cerevisiae* [Kato *et al.*, *Nuc. Acids. Res.*, 22(15): 3104-3112 (1994)], and is hereinafter referred to as *rad3+*. Cells having mutations in *rad3+* fail to either sense or appropriately respond to DNA damage and subsequently lose viability more rapidly than wild type cells after exposure to clastogenic agents or events (*e.g.*, IR, DNA damaging agents,



-3-

and mutations affecting chromosomal integrity). See Weinert *et al.*, *GENES & DEVELOPMENT*, 8: 652-665 (1994) and Al-Khodairy *et al.*, *EMBO J.*, 11(4): 1343-1350 (1992). This sensitivity to IR (radiosensitivity) can be caused by defects in checkpoint responses or defects in direct DNA repair reactions.

5               The product of the *rad3+* gene is an approximately 270 kD protein that falls into a growing family of high molecular weight PIK-related kinases. See Hunter, *Cell*, 83: 1-4 (1995) for a discussion of this family of kinases. The primary structures of the catalytic domains found in members of this gene family are closely related to well characterized phosphatidylinositol kinases. This structural relationship  
10 initially suggested that these PIK-related kinases might be capable of phosphorylating lipids. When the substrate specificity of the PIK-related kinases is examined, however, these enzymes appear to function as protein kinases and have yet to be demonstrated to phosphorylate phosphatidylinositides. Hartley *et al.*, *Cell*, 82: 849-856 (1995) reports that purified preparations highly active in protein kinase assays  
15 failed to show lipid kinase activity. Additional PIK-related kinases identified include: the TEL1 gene product from *S. cerevisiae* which affects telomere length [Greenwell *et al.*, *Cell*, 82: 823-829 (1995)], and *Mei41+* gene product from *Drosophila melanogaster* which is important for a G2 checkpoint and meiotic development [Hari *et al.*, *Cell*, 82: 815-821 (1995)], the DNA-PK gene product from mouse which is  
20 important in immunoglobulin rearrangements and processing of DNA double strand breaks, and the FRAP gene product which is important in the G1/S transition [Brown, E. *et al.*, *Nature*, 377:441-446 (1995)]. Mutations in the DNA-PK gene can result in the Severe Combined Immunodeficiency Syndrome (SCID) defect (Hartley *et al.*, *supra*).

25               In humans, less is known about the molecular components required for checkpoint function. One component of the mammalian checkpoint machinery has been identified through the analysis of the human disease syndrome ataxia-telangiectasia (AT). Patients with AT show a diverse set of clinical symptoms, including predisposition to a variety of tumor types. Fibroblasts from AT patients are  
30 radiosensitive and fail to undergo cell cycle arrest following treatment with IR leading to a phenomenon termed radioresistant DNA synthesis. This is reminiscent of the *S. pombe rad3* defect where cells fail to sense or respond appropriately to DNA damage.

Interestingly, the locus responsible for AT, the Ataxia-Telangiectasia Mutated (ATM) gene, was recently described in Savitsky *et al.*, *Science*, 268: 1749-1753 (1995) and the partial cDNA encodes a protein with amino acid similarity to the *rad3+* gene. Savitsky *et al.*, *Human Molecular Genetics*, 4(11):2025-2032 (1995) describes  
5 isolation of a cDNA encoding full length ATM. The increased radiosensitivity of *rad3+* yeast mutants and of mammalian cells lacking functional ATM suggests that these proteins may comprise a family of checkpoint proteins.

Kuerbitz *et al.*, *Proc. Natl. Acad. Sci. USA*, 89: 7492-7495 (1992) establishes that the tumor suppressor p53 is required for a G1 checkpoint and cell  
10 cycle arrest observed following DNA damage. Irradiation of cells results in increased levels of p53 leading to the transcriptional activation of p53 responsive genes. One such p53-induced target is the product of the WAF1 gene (also called p21, CIP1, and sd1). WAF1 is a member of an expanding class of cell cycle regulators termed cyclin-dependent kinase inhibitory proteins. The activities of cyclin-dependent kinases  
15 control transit through the cell cycle. Transcriptional activation of WAF1 thus provides a direct link between DNA damage-dependent induction of p53 and the inhibition of kinases essential for cell cycle progression. See Elledge and Harper, *Current Opinion in Cell Biology*, 6: 847-852 (1994). Mutations in the p53 gene are one of the most common genetic alterations in human cancers. For example, Baker  
20 *et al.*, *Science*, 244:217-221 (1989) reports that approximately 70% of human colorectal carcinomas contain deletions or mutant copies of the p53 gene. In addition, Fearon *et al.*, *Cell*, 61: 759-767 (1990) reports that breast, lung, bladder and brain tumors have been associated with loss of chromosome 17p, the region to which the p53 gene localizes.

At present there is relatively little known about the molecular  
25 components of the G2 checkpoints in mammalian cells. Caffeine is a chemical entity which abrogates G2 checkpoint control. Russell *et al.*, *Cancer Res.*, 55: 1639-1642 (1995) and Powell *et al.*, *Cancer Res.*, 55: 1643-1648 (1995) report that analysis of cell lines which differ only by the presence or absence of functional p53 demonstrated  
30 preferential caffeine-enhanced sensitization to IR in those cells lacking the p53-dependent G1 checkpoint. Thus, the conversion of potentially lethal damage into

-5-

lethal damage is greater in cells lacking the G1 and G2 checkpoints in comparison to cells containing an intact G1 checkpoint.

While certain cells undergo DNA damage-dependent cell cycle arrest, other cells appear to respond to DNA damage by initiating an intrinsic suicide program termed apoptosis or programmed cell death. The factors determining which process occurs are not fully understood. Recent work has demonstrated an important role for p53 both in the regulation of G1 cell cycle transitions and apoptosis. Symonds *et al.*, *Cell*, 78: 703-711 (1994) describe p53-dependent apoptosis as suppressing tumor growth and progression *in vivo*.

High doses of radiation and chemotherapy are used to treat tumor cells in order to damage DNA so severely that the cells will die. However, even though tumor cells having mutations in the p53 gene are defective in a G1 checkpoint, they can still repair DNA damaged induced by radiation or chemotherapy. The present invention contemplates, for example, that inhibition of the G2 checkpoint in tumor cells should lead to a state in which tumor cells are incapable of repairing DNA damage therefore sensitizing the tumor cells to DNA damaging agents. Normal cells, containing intact G1 and G2 checkpoints, should still be able to repair DNA damage in the presence of a G2 checkpoint-specific inhibitor. Thus, treatment of tumors with a G2 checkpoint-specific inhibitor followed by radiation or chemotherapy should increase the efficacy of cell killing and thereby decrease the required doses of toxic DNA-damaging agents.

There thus exists a need in the art for identification of the mammalian proteins that are involved in the cell cycle checkpoints in order to develop therapies for the human disease states associated with defective cell cycle checkpoints and for the isolation of the genes encoding those proteins which in themselves may be useful as therapeutics or which would enable the development of therapeutically useful modulators of the proteins encoded by the genes.

### SUMMARY OF THE INVENTION

The present invention provides novel human PIK-related kinases essential for a cell cycle checkpoint that responds in the G2 phase of the cell cycle to both damaged and unreplicated DNA.

In one of its aspects, the present invention provides purified and isolated polynucleotides (*e.g.*, DNAs and RNAs, both coding and non-coding strands thereof) encoding the cell cycle checkpoint PIK-related kinase M CCS1 and polynucleotides encoding other cell cycle checkpoint PIK-related kinases that exhibit about 50, about 60, or about 65 % nucleotide identity to the M CCS1 polynucleotide region encoding the M CCS1 kinase domain (M CCS1 $\alpha$  nucleotides 6579 to 7562 of SEQ ID NO: 30 or M CCS1 $\beta$  nucleotides 6457 to 7440 of SEQ ID NO: 32). Alternatively, the M CCS1-like PIK-related kinases exhibit about 40%, about 45 %, or about 50% amino acid identity to the M CCS1 kinase domain (M CCS1 $\alpha$  amino acids 2083 to 2410 of SEQ ID NO: 31 or M CCS1 $\beta$  amino acids 2152 to 2480 of SEQ ID NO: 33). Polynucleotides contemplated by the invention include genomic DNAs, RNAs, cDNAs and wholly or partially chemically synthesized DNAs. Preferred polynucleotides of the invention comprise the M CCS1 $\alpha$  DNA sequence set out in SEQ ID NO: 30, the partial M CCS1 $\beta$  DNA sequence set out in SEQ ID NO: 3, the full length M CCS1 $\beta$  DNA sequence set out in SEQ ID NO: 32, and DNA sequences which hybridize to the noncoding strands thereof under stringent conditions or which would hybridize but for the redundancy of the genetic code. Exemplary stringent hybridization conditions are as follows: hybridization at 65°C in 3X SSC, 20mM NaPO<sub>4</sub> pH 6.8 and washing at 65°C in 0.2X SSC. It is understood by those of skill in the art that variation in these conditions occurs based on the length and GC nucleotide base content of the sequences to be hybridized. Formulas standard in the art are appropriate for determining exact hybridization conditions. See Sambrook *et al.*, 9.47-9.51 in *Molecular Cloning*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1989). The M CCS1 $\alpha$  DNA of SEQ ID NO: 30 was deposited with the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, Maryland 20852, on November 3, 1995 as an insert in plasmid pBSHFB/HT2-27 in *E. coli* DH5 $\alpha$  and was assigned ATCC Accession No. 69951. The M CCS1 $\beta$  DNA of SEQ ID NO: 32, was deposited with the ATCC on November 7, 1995 as an insert in plasmid 517 in *E. coli* DH5 $\alpha$  and was assigned ATCC Accession No. 69950.

The DNA sequence information provided by the present invention makes possible the identification and isolation of DNAs encoding related molecules

-7-

by well-known techniques such as DNA/DNA hybridization as described above and polymerase chain reaction (PCR) cloning. As one series of examples, knowledge of the sequence of a cDNA encoding MCCI1 makes possible the isolation by DNA/DNA hybridization of genomic DNA sequences encoding the kinase and expression control regulatory sequences such as promoters, operators and the like. Similarly, knowledge of a partial cDNA sequence encoding MCCI1 $\beta$  make isolation of a complete cDNA possible. DNA/DNA hybridization procedures carried out with DNA sequences of the invention under stringent conditions are likewise expected to allow the isolation of DNAs encoding allelic variants of the PIK-related kinase; non-human species enzymes homologous to the PIK-related kinase; and other structurally related proteins sharing one or more of the enzymatic activities, or abilities to interact with members or regulators, of the cell cycle checkpoint pathway in which MCCI1 participates. Polynucleotides of the invention when detectably labelled are also useful in hybridization assays to detect the capacity of cells to synthesize MCCI1. The DNA sequence information provided by the present invention also makes possible the development, by homologous recombination or "knockout" strategies [see, Capecchi, *Science*, 244: 1288-1292 (1989)], of rodents that fail to express functional MCCI1 or that express a variant of MCCI1. Such rodents are useful as models for studying the activities of MCCI1 and MCCI1 modulators *in vivo*. Polynucleotides of the invention may also be the basis for diagnostic methods useful for identifying a genetic alteration(s) in the MCCI1 locus that underlies a disease state or states. Also made available by the invention are anti-sense polynucleotides relevant to regulating expression of MCCI1 by those cells which ordinarily express the same.

The invention also provides autonomously replicating recombinant constructions such as plasmid and viral DNA vectors incorporating polynucleotides of the invention, especially vectors in which the polynucleotides are functionally linked to an endogenous or heterologous expression control DNA sequence and a transcription terminator.

According to another aspect of the invention, host cells, especially unicellular host cells such as procaryotic and eukaryotic cells, are stably transformed or transfected with DNAs of the invention in a manner allowing expression of the PIK-related kinase therein. Host cells of the invention are conspicuously useful in

methods for the large scale production of MCCS1 wherein the cells are grown in a suitable culture medium and the desired enzymes are isolated from the cells or from the medium in which the cells are grown.

5 MCCS1 products having part or all of the amino acid sequence set out in SEQ ID NO: 31, SEQ ID NO: 4, or SEQ ID NO: 33 are contemplated. Use of mammalian host cells is expected to provide for such post-translational modifications (*e.g.*, myristoylation, glycosylation, truncation, lipidation and tyrosine, serine or threonine phosphorylation) as may be needed to confer optimal biological activity on recombinant expression products of the invention. The enzyme products of the invention may be full length polypeptides, fragments or variants. Variants comprise 10 MCCS1 products wherein one or more of the specified (*i.e.*, naturally encoded) amino acids is deleted or replaced or wherein one or more nonspecified amino acids are added: (1) without loss of the kinase activity specific to MCCS1; or (2) with disablement of the kinase activity specific to MCCS1; or (3) with disablement of the ability to interact with members or regulators of the cell cycle checkpoint pathway. 15 Substrates of MCCS1 and proteins which interact with MCCS1 may be identified by various assays.

Substrates of MCCS1 may be identified by incorporating test compounds in assays for kinase activity. MCCS1 kinase is resuspended in kinase 20 buffer and incubated either in the presence or absence of the test compound (*e.g.*, casein, histone H1, or appropriate substrate peptide). Moles of phosphate transferred by the kinase to the test compound are measured by autoradiography or scintillation counting. Transfer of phosphate to the test compound is indicative that the test compound is a substrate of the kinase.

25 Interacting proteins may be identified by the following assays.

A first assay contemplated by the invention is a two-hybrid screen. The two-hybrid system was developed in yeast [Chien *et al.*, *Proc. Natl. Acad. Sci. USA*, 88: 9578-9582 (1991)] and is based on functional *in vivo* reconstitution of a transcription factor which activates a reporter gene. Specifically, a polynucleotide 30 encoding a protein that interacts with MCCS1 is isolated by: transforming or transfecting appropriate host cells with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA

-9-

binding domain and an activating domain; expressing in the host cells a first hybrid DNA sequence encoding a first fusion of part or all of MCCS1 and either the DNA binding domain or the activating domain of the transcription factor; expressing in the host cells a library of second hybrid DNA sequences encoding second fusions of part or all of putative MCCS1 binding proteins and the DNA binding domain or activating domain of the transcription factor which is not incorporated in the first fusion; detecting binding of an MCCS1 interacting protein to MCCS1 in a particular host cell by detecting the production of reporter gene product in the host cell; and isolating second hybrid DNA sequences encoding the interacting protein from the particular host cell. Presently preferred for use in the assay are a *lexA* promoter to drive expression of the reporter gene, the *lacZ* reporter gene, a transcription factor comprising the *lexA* DNA binding domain and the GAL4 transactivation domain, and yeast host cells.

Other assays for identifying proteins that interact with MCCS1 may involve immobilizing MCCS1 or a test protein, detectably labelling the nonimmobilized binding partner, incubating the binding partners together and determining the amount of label bound. Bound label indicates that the test protein interacts with MCCS1.

Another type of assay for identifying MCCS1 interacting proteins involves immobilizing MCCS1 or a fragment thereof on a solid support coated (or impregnated with) a fluorescent agent, labelling a test protein with a compound capable of exciting the fluorescent agent, contacting the immobilized MCCS1 with the labelled test protein, detecting light emission by the fluorescent agent, and identifying interacting proteins as test proteins which result in the emission of light by the fluorescent agent. Alternatively, the putative interacting protein may be immobilized and MCCS1 may be labelled in the assay.

Also comprehended by the present invention are antibody products (e.g., monoclonal and polyclonal antibodies, single chain antibodies, chimeric antibodies, CDR-grafted antibodies and the like) and other binding proteins (such as those identified in the assays above) which are specific for the MCCS1 kinases of the invention. Binding proteins can be developed using isolated natural or recombinant enzymes. The binding proteins are useful, in turn, for purifying recombinant and

-10-

naturally occurring enzymes and identifying cells producing such enzymes. Specifically illustrating monoclonal antibodies of the invention are the monoclonal antibodies produced by hybridoma cell lines 224C and 224F which were deposited with the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, MD 20852 on November 7, 1996 and assigned ATCC Accession Nos. HB 12233 and HB 12234, respectively. Assays for the detection and quantification of proteins in cells and in fluids may involve a single antibody substance or multiple antibody substances in a "sandwich" assay format. The binding proteins are also manifestly useful in modulating (*i.e.*, blocking, inhibiting, or stimulating) enzyme/substrate or enzyme/regulator interactions. Anti-idiotypic antibodies specific for PIK-related kinase binding proteins are also contemplated.

The invention contemplates that mutations in the M CCS1 gene that result in loss of normal function of the M CCS1 gene product underlie human disease states in which failure of the G<sub>2</sub> cell cycle checkpoint is involved. Gene therapy to restore M CCS1 activity would thus be indicated in treating those disease states (for example, testicular cancer). Delivery of a functional M CCS1 gene to appropriate cells is effected *in vivo* or *ex vivo* by use of viral vectors (*e.g.*, adenovirus, adeno-associated virus, or a retrovirus) or *ex vivo* by use of physical DNA transfer methods (*e.g.*, liposomes or chemical treatments). For reviews of gene therapy technology see Friedmann, *Science*, 244: 1275-1281 (1989); Verma, *Scientific American*: 68-84 (1990); and Miller, *Nature*, 357: 455-460 (1992). Alternatively, it is contemplated that in other human disease states preventing the expression of or inhibiting the activity of M CCS1 will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate the expression of M CCS1. Antisense nucleic acids (preferably 10 to 20 base pair oligonucleotides) capable of specifically binding to M CCS1 expression control sequences or M CCS1 RNA are introduced into cells (*e.g.*, by a viral vector or colloidal dispersion system such as a liposome). The antisense nucleic acid binds to the M CCS1 target sequence in the cell and prevents transcription or translation of the target sequence. Phosphothioate and methylphosphate antisense oligonucleotides are specifically contemplated for therapeutic use by the invention. The antisense



-11-

oligonucleotides may be further modified by poly-L-lysine, transferrin polylysine, or cholesterol moieties at their 5' end.

Moreover, for example, if a particular form of cancer results from a mutation in a gene other than M CCS1 such as the p53 gene, an agent which inhibits the transcription or the enzymatic activity of M CCS1 and thus the G<sub>2</sub> cell cycle checkpoint may be used to render cancerous cells more sensitive to chemotherapy or radiation therapy. The therapeutic value of such an agent lies in the fact that current radiation therapy or chemotherapy in most cases does nothing to overcome the ability of the p53 mutant cancerous cell to sense and correct the DNA damage imposed as a result of the treatment. As a result, a cancer cell can simply repair the DNA damage. Modulating agents of the invention may therefore be chemotherapy and radiation adjuvants or may be directly active as chemotherapeutic drugs themselves.

Agents that modulate M CCS1 kinase activity may be identified by incubating a test compound with M CCS1 immunopurified from cells naturally expressing the PIK-related kinase, with M CCS1 obtained from recombinant procaryotic or eukaryotic host cells expressing the enzyme, or with purified M CCS1, and then determining the effect of the test compound on M CCS1 activity. The activity of the PIK-related kinase can be measured by determining the moles of <sup>32</sup>P-phosphate transferred by the kinase from gamma-<sup>32</sup>P-ATP to either itself (autophosphorylation) or to an exogenous substrate such as a lipid or protein. The amount of phosphate incorporated into the substrate is measured by scintillation counting or autoradiography. An increase in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the test compound is an activator of said M CCS1 kinase. Conversely, a decrease in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the modulator is an inhibitor of said M CCS1 kinase. In another aspect, agents that modulate both M CCS1 and ATM or modulate one of the enzymes are also contemplated. Agents which modulate M CCS1 are screened in a kinase assay as described above in which ATM is the phosphorylating enzyme.

-12-

In a presently preferred assay, a MCCC1-specific antibody linked to agarose beads is incubated with a cell lysate prepared from host cells expressing the kinase. The beads are washed to remove proteins binding nonspecifically to the beads and the beads are then resuspended in kinase buffer. The reaction is initiated by the addition of gamma-<sup>32</sup>P-ATP and an appropriate exogenous substrate such as lipid or peptide. The activity of the kinase is measured by determining the moles of <sup>32</sup>P-phosphate transferred either to the kinase itself or the added substrate. In a preferred embodiment the host cells lack endogenous MCCC1 and/or ATM kinase activity. The selectivity of a compound that modulates the kinase activity of MCCC1 can be evaluated by comparing its activity on MCCC1 to its activity on other known PIK-related kinases. The combination of the recombinant MCCC1 products of the invention with other recombinant PIK-related kinase products in a series of independent assays provides a system for developing selective modulators of MCCC1.

Furthermore, combinatorial libraries, peptide and peptide mimetics, defined chemical entities, oligonucleotides, and natural product libraries may be screened for activity as modulators in assays such as those described below.

For example, an assay for identifying modulators of MCCC1 kinase activity involves incubating an MCCC1 kinase preparation in kinase buffer with gamma-<sup>32</sup>P-ATP and an exogenous kinase substrate, both in the presence and absence of a test compound, and measuring the moles of phosphate transferred to the substrate. An increase in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the test compound is an activator of said MCCC1 kinase. Conversely, a decrease in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the modulator is an inhibitor of said MCCC1 kinase.

Moreover, assays for identifying compounds that modulate interaction of MCCC1 with other proteins may involve: transforming or transfecting appropriate host cells with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA-binding domain and an activating domain; expressing in the host cells a first hybrid DNA sequence encoding

-13-

a first fusion of part or all of M CCS1 and the DNA binding domain or the activating domain of the transcription factor; expressing in the host cells a second hybrid DNA sequence encoding part or all of a protein that interacts with M CCS1 and the DNA binding domain or activating domain of the transcription factor which is not incorporated in the first fusion; evaluating the effect of a test compound on the interaction between M CCS1 and the interacting protein by detecting binding of the interacting protein to M CCS1 in a particular host cell by measuring the production of reporter gene product in the host cell in the presence or absence of the test compound; and identifying modulating compounds as those test compounds altering production of the reported gene product in comparison to production of the reporter gene product in the absence of the modulating compound. Presently preferred for use in the assay are a *lexA* promoter to drive expression of the reporter gene, the *lacZ* reporter gene, a transcription factor comprising the *lexA* DNA binding domain and the GAL4 transactivation domain, and yeast host cells.

Another type of assay for identifying compounds that modulate the interaction between M CCS1 and an interacting protein involves immobilizing M CCS1 or a natural M CCS1 interacting protein, detectably labelling the nonimmobilized binding partner, incubating the binding partners together and determining the effect of a test compound on the amount of label bound wherein a reduction in the label bound in the presence of the test compound compared to the amount of label bound in the absence of the test compound indicates that the test agent is an inhibitor of M CCS1 interaction with protein. Conversely, an increase in the bound in the presence of the test compound compared to the amount label bound in the absence of the compound indicates that the putative modulator is an activator of M CCS1 interaction with the protein.

Yet another method contemplated by the invention for identifying compounds that modulate the binding between M CCS1 and an interacting protein involves immobilizing M CCS1 or a fragment thereof on a solid support coated (or impregnated with) a fluorescent agent, labelling the interacting protein with a compound capable of exciting the fluorescent agent, contacting the immobilized M CCS1 with the labelled interacting protein in the presence and absence of a test compound, detecting light emission by the fluorescent agent, and identifying

-14-

modulating compounds as those test compounds that affect the emission of light by the fluorescent agent in comparison to the emission of light by the fluorescent agent in the absence of the test compound. Alternatively, the M CCS1 interacting protein may be immobilized and M CCS1 may be labelled in the assay.

5           The present invention further provides a cell-based complementation assay for identifying compounds which modulate the activity of M CCS1 or ATM. The assay involves complementation of a phenotypic trait associated with a genetic alteration in the cell. For example, the genetic alteration identified as *esr1-1* results in cellular sensitivity to DNA damage in yeast cells [Kato *et al.*, *Nuc. Acids. Res.*,  
10   22(15): 3104-3112 (1994)]. *esr1-1* cells fail to either sense or appropriately response to DNA damage after exposure to DNA damaging agents such as ionizing radiation or clastogenic agents. The phenotypic trait of the genetically altered cell is complemented by transforming and expressing M CCS1 or ATM in the cell. The transformed cells are exposed to DNA damaging treatment (e.g. ionizing radiation)  
15 in the presence and absence of a test compound and sensitivity of the cells to DNA damage is measured. Agents that affect the cell sensitivity to DNA damaging activity of M CCS1 and/or ATM are identified as modulators.

          Modulators of M CCS1 may affect its kinase activity, its localization in the cell, and/or its interaction with members of the cell cycle checkpoint pathway.  
20 M CCS1 modulators may be formulated in compositions comprising pharmaceutically acceptable carriers. Such compositions may additionally include chemotherapeutic agents. Dosage amounts indicated would be sufficient to result in modulation of M CCS1 activity *in vivo*. Selective modulators may include, for example, polypeptides or peptides which specifically bind to M CCS1 or M CCS1 nucleic acid,  
25 oligonucleotides which specifically bind to the PIK-related kinase or PIK-related kinase nucleic acid, and/or other non-peptide compounds (e.g., isolated or synthetic organic molecules) which specifically react with M CCS1 or M CCS1 nucleic acid. Mutant forms of M CCS1 which affect the enzymatic activity or cellular localization of wild-type M CCS1 are also contemplated by the invention. Presently preferred  
30 regions of the PIK-related kinases which are targets for the development of selective modulators include, for example, the following four regions: the M CCS1 $\alpha$  amino terminal effector domain (amino acids 1 to 1081 of SEQ ID NO: 31), the M CCS1 $\beta$

-15-

amino terminal effector domain (amino acids 1 to 1150 of SEQ ID NO: 33), the  
MCCS1 $\alpha$  *rad3*+ domain (amino acids 1082 to 2082 of SEQ ID NO: 31), the  
MCCS1 $\beta$  *rad3*+ domain (amino acids 1151 to 2151 of SEQ ID NO: 33), the  
MCCS1 $\alpha$  PIK domain (amino acids 2083 to 2410 of SEQ ID NO: 31), and the  
5 MCCS1 $\beta$  PIK domain (amino acids 2152 to 2480 of SEQ ID NO: 33).

### DETAILED DESCRIPTION

The present invention is illustrated by the following examples. Example 1 details the isolation of cDNAs encoding M CCS1 kinases. Example 2 describes mapping of the human M CCS1 gene to human chromosome 3. The recombinant expression of M CCS1 in *E. coli* and insect cells is respectively described in Examples 3 and 4. Example 4 also presents assays for measuring M CCS1 kinase activity. Example 5 describes the production of M CCS1-specific polyclonal and monoclonal antibodies. Example 6 reports the immunoprecipitation of M CCS1 kinase associated activity from mouse testes. Example 7 examines the expression of M CCS1 mRNA in various human tissues and cancer cell lines. Example 8 describes analyses of M CCS1 mRNA and protein expression in mouse testes. Example 9 describes analyses of M CCS1 protein expression in meiotic cells. Assays for substrates and interacting proteins of M CCS1 are described in Example 10. Example 11 describes modulators and assays for modulators of the kinase activity of M CCS1. Example 12 describes the cell-based complementation assay for identifying modulators of M CCS1 and/or ATM and Example 13 describes the kinase activity of ATM.

#### Example 1

cDNAs encoding the PIK-related kinase M CCS1 were isolated by a series of PCR reactions.

An alignment of the amino acid sequences of *S. pombe rad3+* (Hari *et al.*, *supra*) and *S. cerevisiae MEC1* (Kato *et al.*, *supra*) was the basis for design of seven degenerate oligonucleotides that encoded (or were complementary to) the regions of highest homology/lowest degeneracy between the sequences and contained convenient restriction sites to facilitate cloning of amplification products. The oligonucleotides were then used in a PCR-based assay to isolate a related human sequence.

Initially, PCR amplifications were performed on cDNA preparations from rat T-cells, human peripheral blood mononuclear cells (PBMC), and *S. cerevisiae* genomic DNA. Five oligonucleotide pairs were used (oDH15a/oDH16, oDH15b/oDH16, oDH17a/oDH16, oDH15a/oDH17b, and oDH15b/oDH17b) for the primary amplifications. The sequences of the oligonucleotide primers included

-17-

inosines and are set out below in IUPAC nomenclature for degenerate nucleotide positions.

oDH15a (SEQ ID NO: 5)

5' GCA GAC GGA TCC GGI WCI GAY GGI AAY HTI TAY 3'

5

oDH15b (SEQ ID NO: 6)

5' GCA GAC GGA TCC GGI WCI GAY GGI AAY 3'

oDH16 (SEQ ID NO: 7)

5' GCA GAC GAA TTC RCA RTY RAA RTC IAC RTG 3'

10

oDH17a (SEQ ID NO: 8)

5' GCA GAC GGA TCC AAR TTY

CCI CCI RTI YTI TAY SAR TGG TT 3'

oDH17b (SEQ ID NO: 9)

5' GCA GAC GAA TCC AAC CAY

TSR TAI ARI AYI GGI GGR AAY TT 3'

15 PCR was performed on reaction mixtures of 1X PCR buffer (Perkin Elmer Cetus, Emeryville, California), 2-3 $\mu$ M oDH primers, 1.5mM MgCl<sub>2</sub>, 200 $\mu$ M dNTPs, and 0.5  $\mu$ l Amplitaq polymerase. The reaction was performed in a Perkin-Elmer Cetus Thermocycler Model 480 under the following conditions: denaturation at 94°C for 1 minute, annealing at 64°C for 2 minutes, and elongation at 72°C for 1 minute for 3 cycles. The procedure was then repeated using 60°C annealing temperature for 3 cycles, 56°C annealing for 3 cycles, and finished with denaturation at 94°C for 1 minute, annealing at 54°C (2 minutes, and elongation at 72°C for 1 minute for 30 cycles. PCR products were separated on 2 or 4% Tris Acetate EDTA (TAE) agarose gels, stained with ethidium bromide, and DNA products were visualized by UV  
25 fluorescence.

-18-

From the primary amplifications of yeast genomic DNA, rat T-cell cDNA, and human PBMC cDNA, only a single reaction with yeast genomic DNA (oDH17a/oDH16) gave a visible amplification product, resulting in a product that was the expected size for the region of the *S. cerevisiae MEC1* gene between these primers. Further analysis of the oDH17a/oDH16 amplifications that utilized rat T-cell and PBMC cDNA was therefore performed. To remove oligonucleotides and "primer dimers" that might interfere with subsequent PCR, primary reactions were purified prior to reamplification.

A "nested" PCR strategy was employed, and amplifications were repeated with primer pairs oDH18a/oDH16 and oDH18b/oDH16 under reaction conditions described above with cycle times of denaturation of 94°C for 1 minute, annealing at 55°C for 1 minute, and elongation at 72°C for 30 seconds for 30 cycles. The sequences of the oDH18a and oDH18b oligonucleotide primers included inosines and are set out below in IUPAC nomenclature for degenerate nucleotide positions.

oDH18a (SEQ ID NO: 10)

5' GCA GAC GGA TCC YTI GGI YTI GGI GAY CGI CA 3'

oDH18b (SEQ ID NO: 11)

5' GCA GAC GGA TCC YTI GGI YTI GGI GAY AGR CA 3'

An approximately 90 base pair (bp) product (the expected size amplification product for these primers) was seen in the reamplifications of the yeast genomic and human PBMC cDNA primary reactions. No 90 bp product was seen in the reamplification of the primary reaction on rat T-cell cDNA and this reaction was not analyzed further.

In addition to the approximately 90 bp product, several other non-specific bands were also present, though significantly fewer than were observed when the primary reactions were reamplified with oDH17a/oDH16. While the approximately 90 bp product was present in both the oDH18a/oDH16 and oDH18b/oDH16 reamplifications of the yeast genomic DNA primary reactions, only the oDH18a/oDH16 reaction yielded the appropriate size fragment during



-19-

reamplification of the human PMBC cDNA primary reaction. This was presumed to reflect codon usage in the human gene (compare primers oDH18a and oDH18b). The approximately 90 bp product from the oDH18a/oDH16 reamplification of the human PMBC cDNA primary reaction was gel purified and subcloned into the pBluescript SKII+ cloning vector (Stratagene, La Jolla, California) and sequenced.

Analysis of the sequence encoded by the 90 bp product indicated that the deduced amino acid sequence was similar to both *S. cerevisiae* MEC1 and *S. pombe rad3+*, but was not identical to either. To identify a larger region of coding sequence and extend the sequence comparison, a non-degenerate oligonucleotide, oDH23 5' GACGCAGAATTCACCAAGTCAAAGAATCAAAGAG 3' (SEQ ID NO: 12), was synthesized for use in additional amplification reactions. Reamplification of the purified PBMC cDNA primary reaction with oDH17a/oDH23 led to the production of an amplification product of 174 bp. This fragment was then purified, subcloned and sequenced as described above. Computer analysis of the conceptual translation product confirmed its relationship (similar but not identical) to *MEC1* and *rad3+*. This PCR fragment was then used as a probe to screen a plasmid library containing macrophage cDNA using the following hybridization conditions: incubation of nitrocellulose filters with radiolabelled probes in 3X SSC, 5X Denhardt's, 0.1% sarcosyl, 20mM NaPO<sub>4</sub> pH 6.8, 100 ug/ml single stranded salmon sperm DNA, for 18 to 24 hours at 65°C. Washes were done 3 times in 0.2X SSC, 0.1% SDS at 65°C for 30 minutes (with changes of wash buffer). Four positive clones were isolated, and the nucleotide sequence of each was determined. Computer analysis of the four sequences demonstrated that they were overlapping clones derived from a locus with homology to the *rad3+* gene from *S. pombe*. Clone 517 (ATCC 69950) contained a 2.8 kbp insert and its DNA and deduced amino acid sequence are set out in SEQ ID NOs: 3 and 4, respectively. The clone contained an open reading frame encoding an amino terminal truncated protein product of 870 amino acids which were 39% identical to the COOH-terminus of *rad3+*. The protein product of the cDNA insert was named MCCS1β.

The sequence of clone 517 was used to design the oligonucleotides, mo3 5'-CTACAGAGCCAAGGAG-3' (SEQ ID NO: 13) and mo6 5'-TCGAGCTATGCTACTAGTGGGC-3' (SEQ ID NO: 14), which were used to

-20-

generate a probe using a gel purified EcoRI fragment derived from clone 517 as a template. The PCR conditions were as follows: 50 ng DNA fragment, 1X PCR buffer (Perkin-Elmer Cetus), 1.5mM MgCl<sub>2</sub>, 200μM dATP, dGTP, and TTP, 1μM dCTP, 50μCi α<sup>32</sup>P-dCTP, 10ng/ml each oligonucleotide, 1U AmpliTaq (Perkin-Elmer Cetus). The reaction was performed in a Perkin-Elmer Cetus Thermocycler Model 480 for an initial denaturing cycle at 94°C for 4 minutes followed by 20 cycles of 94°C for 15 seconds, 60°C for 15 seconds, 72°C for 30 seconds. Unincorporated nucleotides were removed using a Stratagene Nuc-trap Push Column.

Since Northern blot analyses showed that the expression of the mRNA corresponding to clone 517 was highest in testis, one million clones from a human testis cDNA library (Stratagene #939202) were screened with the PCR-generated probe and eleven clones were obtained. The two longest clones, HT2 and HT9, were chosen for analysis. HT2 contained a 4.7 Kb insert (corresponding to nucleotide 2974 of SEQ ID NO: 30 and extending further downstream than SEQ ID NO: 1) and HT9 contained a 5485 bp insert (corresponding to nucleotides 2152 to 7624 of SEQ ID NO: 30). Nucleotide sequence analysis revealed that in the region common to both cDNA clones there was a single base pair insertion of a T at nucleotide 3233 in HT9. This nucleotide insertion causes the predicted amino acid reading frame to shift and then terminate and is believed to be an error introduced by reverse transcriptase in clone HT9.

In order to isolate a clone containing an additional 2.5 Kb, one million clones from each of three additional cDNA libraries were screened: a human fetal brain cDNA library (Stratagene #93206), a human heart cDNA library (Stratagene #936207), and a human aorta cDNA library (Clontech Laboratories #HL1136a, Palo Alto, California). The sequence of the most 5' region of HT9 was utilized to design and synthesize two oligonucleotides, oHT9-1 5'-CCTAGTCCAGTAAACTTGC-3' (SEQ ID NO: 15) and oHT9-4 5'-TTTGC GGCCCTTCCAATATC-3' (SEQ ID NO: 16) which were used to generate a 317 bp PCR probe under conditions described for generating the probe above. While no positive clones were isolated from the heart or aorta cDNA libraries, two positive clones were obtained from the fetal brain library. One of these clones, HFB2, included a cDNA 4.5 Kb insert which included

-21-

approximately 2300 bp of additional sequence. The HFB2 insert corresponds to nucleotides 1 to 3194 of SEQ ID NO: 30.

A composite cDNA encoding  $\text{MCCS1}\alpha$  was constructed from clones HFB2, HT9 and HT2. The three clones were joined together by digesting HFB2 with the restriction enzymes KpnI and Sall to generate a fragment to comprise the 5' end of the composite clone, digesting HT9 with KpnI and NotI to generate a fragment to comprise the 3' end of the composite clone, and then ligating isolated fragments to the vector pBS SK<sup>+</sup> (Stratagene) that had been digested with Sall and NotI. The region of the HT9 fragment containing the one nucleotide insertion was replaced with an EcoRV fragment containing nucleotides 3174 to 5282 of clone HT2. The final plasmid containing a 7621 bp insert was named pBSHFB2HT2-27 (ATCC 69951). The DNA and deduced amino acid sequence of the insert are presented in SEQ ID NOs: 1 and 2, respectively. The coding domain of the cDNA initiates with an ATG at nucleotide 333 and ends with a termination codon at nucleotide 7560 predicting a coding sequence of 2409 amino acids and protein of 265 kD. The protein product of the cDNA insert was named  $\text{MCCS1}\alpha$ . Subsequent sequence analysis of the insert in plasmid pBSHFB2HT2-27 (ATCC 69951) revealed sequencing errors in SEQ ID NO: 1. Corrected DNA and deduced amino acid sequences of the insert are set out in SEQ ID NOs: 23 and 24, respectively. Even further sequence analysis of insert in plasmid pBSHFB2HT2-27 revealed sequencing error in SEQ ID NO: 23. At nucleotide position 6317 (SEQ ID NO: 23) a "G" was erroneously included and between positions 6338 and 6339 the sequence was missing an "A". The corrected sequences of  $\text{MCCS1}\alpha$  are provided in SEQ ID NOs: 30 and 31.

Comparison of the predicted amino acid sequence of  $\text{MCCS1}\alpha$  with the partial amino acid sequence of  $\text{MCCS1}\beta$  predicted from clone 517 revealed the presence of a seventy amino acid deletion in the  $\text{MCCS1}\alpha$  product. The  $\text{MCCS1}\beta$  clone 517 amino acid sequence corresponds to  $\text{MCCS1}\alpha$  amino acids 1611 to 2410 of SEQ ID NO: 31. The seventy amino acid deletion in  $\text{MCCS1}\alpha$  (*i.e.*, where the seventy amino acids would be inserted to generate a product identical to  $\text{MCCS1}\beta$ ) occurs between amino acids 2065 and 2066 in SEQ ID NO: 31, seventeen amino acids upstream from the kinase domain. Since both clones maintain an open reading frame, cDNA clone pBSHFB2HT2-27 was apparently generated from alternatively

-22-

spliced mRNA. The carboxyl terminal domains containing the kinase domains are identical in MCCS1 $\alpha$  (amino acids 2083 to 2410 of SEQ ID NO: 31) and MCCS1 $\beta$  clone (amino acids 543 to 870 of SEQ ID NO: 4).

5 A composite clone containing the complete coding sequence of MCCS1 $\beta$  (with the seventy amino acid insert) is presented in SEQ ID NO: 32. The amino acid sequence deduced from the clone is presented in SEQ ID NO: 33. This clone is constructed by replacing the sequence between the BSTXI site, which cleaves after nucleotide 3229, and the NotI site in the polylinker sequence at the 3' end of pBSHFB2HT2-27 (SEQ ID NO: 1) with the sequence contained in HT2 between the  
10 BstXI site and the NotI site at the 3' end of HT2. Thus this clone contains sequences that are identical to MCCS1 $\alpha$  nucleotides 1 to 5159 of SEQ ID NO: 1 (encoding amino acids 1 to 1609 of SEQ ID NO: 2) linked to sequences that are identical to clone 517 nucleotides 1 to 2610 of SEQ ID NO: 3 (encoding amino acids 1 to 870 of SEQ ID NO: 4). As noted above, subsequent sequence analysis revealed errors  
15 in nucleotides 1 to 5159 of SEQ ID NO: 1. Corrected MCCS1 $\beta$  DNA and deduced amino acid sequences that include the same corrections that appear in MCCS1 $\alpha$  SEQ ID NOs: 23 and 24 are set out in SEQ ID NOs: 25 and 26. The SEQ ID NO: 25 clone represents a cDNA encoding a full length MCCS1 $\beta$  kinase. Further sequences for MCCS1 $\beta$  including corrections of errors identified in resequencing the MCCS1 $\alpha$  clone are presented in SEQ ID NOs: 32 and 33.  
20

The MCCS1 products can be divided into three regions based on similarity to other PIK-related kinases: an amino terminal domain (MCCS1 $\alpha$  amino acids 1 to 1081 of SEQ ID NO: 31 and MCCS1 $\beta$  amino acids 1 to 1150 of SEQ ID NO: 33), a region with similarity to *rad3+* (MCCS1 $\alpha$  amino acids 1082 to 2082 of  
25 SEQ ID NO: 31 and MCCS1 $\beta$  amino acids 1151 to 2151 of SEQ ID NO: 33) and a PIK domain (MCCS1 $\alpha$  amino acids 2083 to 2410 of SEQ ID NO: 31 and MCCS1 $\beta$  amino acids 2152 to 2480 of SEQ ID NO: 33) including a kinase domain. The amino terminal region and *rad3+* region are regulatory domains that modulate the kinase activity of the enzyme and are involved in interactions with associated proteins.  
30

Results of comparisons of the nucleotide and amino acid sequence of MCCS1 $\alpha$  and MCCS1 $\beta$  to the sequences of other PIK-related and non-PIK-related kinases are shown in Table 1. Specifically, the 3' end of MCCS1 $\alpha$  (nucleotides 6579

-23-

to 7562 of SEQ ID NO: 30 encoding the kinase domain), the 3' end of MCCS1 $\beta$  (nucleotides 1627 to 2379 of SEQ ID NO: 32 encoding the kinase domain), the *rad3+* domain of MCCS1 $\alpha$  (nucleotides 3576 to 6578 of SEQ ID NO: 30), and the *rad3+* domain of MCCS1 $\beta$  (clone 517 nucleotides 1 to 1626 of SEQ ID NO: 3) were compared to the analogous region in human ATM [Savitsky *et al.*, *supra*], human DNA-PK [Huntley *et al.*, *Cell*, 82: 849-856 (1995)], human FRAP [Brown *et al.*, *supra*], human p110 [Hu *et al.*, *Mol. Cell. Biol.*, 13(12): 7677-7688 (1993)], *S. cerevisiae* MEC1 [Weinert *et al.*, *Genes Dev.*, 8(6): 652-665 (1994)], *S. pombe* *rad3+* [Seaton *et al.*, *supra* and Hari *et al.*, *Cell*, 82: 815-821 (1995)] and an cAMP-dependent protein kinase (PKA) [Beebe *et al.*, *Mol. Endocrinol.*, 4(3): 465-475 (1990)]. Percent identity of nucleotides is shown in the top line, percent identity of amino acids is shown in the middle line, and percent similarity of amino acids (*i.e.*, including identical amino acids and conservative variations in amino acids) is shown in the bottom line for each kinase in Table 1. Conservative variation as used herein denotes biologically similar residues. Examples of conservative variations include the substitution of one hydrophobic residue such as isoleucine, valine, leucine or methionine for another, or the substitution of one polar residue for another, such as the substitution of arginine for lysine, glutamic for aspartic acids, or glutamine for asparagine, and the like. In the Table, "ND" indicates a value was not determined either because the nucleotide sequence encoding the kinase (*i.e.*, *rad3+*) was not publically available or because the kinase (*i.e.*, FRAP, p110 $\beta$ , or PKA) lacks the particular domain being compared.

-24-

Table 1

	Protein Kinase	MCCS1 $\alpha$ /MCCS1 $\beta$ Kinase Domain	MCCS1 $\alpha$	MCCS1 $\beta$
			<i>rad3</i> + Domain	<i>rad3</i> + Domain
5	<i>S. pombe rad3</i> +	ND	ND	ND
		56	22	30
		72	46	53
	<i>S. cerevisiae</i> <i>MEC1</i>	51	42	44
		45	21	24
		63	46	49
	Human ATM	50	41	41
		38	22	24
		60	46	47
	Human DNA-PK	43	39	43
		29	19	20
		53	45	49
10	Human FRAP	45	ND	ND
		37	ND	ND
		61	ND	ND
	Human p110 $\beta$	45	ND	ND
		24	ND	ND
		54	ND	ND
	Human PKA	39	ND	ND
		16	ND	ND
		39	ND	ND

-25-

**Example 2**

The MCCS1 gene was mapped to chromosome 3 by a PCR-based assay. Human/rodent somatic cell hybrids containing various human chromosome panels available from the NIGMS Human Genetic Mutant Cell Repository [Drwinga  
5 *et al.*, *Genomics*, 16: 311-314 (1993)] were used as templates.

Two oligonucleotide primers oDH23 (SEQ ID NO: 12) and oDH26 5' TGGTTTCTGAGAACATTCCCTGA 3' (SEQ ID NO: 19) based on the MCCS1 $\alpha$  cDNA sequence were utilized to amplify a portion of the gene. The primers generate 237 bp PCR products. PCR conditions consisted of 50 ng genomic DNA, 0.5  $\mu$ g of  
10 each primer, 200  $\mu$ M dNTPs, 1.5mM MgCl<sub>2</sub>, 1X PCR buffer (Perkin Elmer-Cetus), and 1 unit of Amplitaq polymerase (Perkin-Elmer Cetus) in a 25  $\mu$ l reaction volume. The samples were denatured for 4 minutes and then cycled 35 times with denaturing, annealing, and extension times of 45 seconds, 30 seconds, and 45 seconds, respectively, in a Model 480 Cetus Thermocycler. Five  $\mu$ l of the resulting PCR  
15 product was electrophoresed on a 3% agarose gel and stained with ethidium bromide. DNA corresponding to the human/rodent chromosome 3 hybrid yielded a positive amplification product.

In a second set of amplification reactions, the same oligonucleotide primers were used to sublocalize the MCCS1 gene to a specific region on  
20 chromosome 3. The templates for these amplifications consisted of DNA samples from patients with chromosome 3 truncations [Leach *et al.*, *Genomics*, 24: 549-556 (1994)]. Amplifications were performed as described in the foregoing paragraph. The pattern of positive amplification products narrowed the localization to the interval between q21 and q25.1.

**Example 3**

Polynucleotides encoding carboxyl terminal portions of the PIK-related kinase MCCS1 $\beta$  were expressed by recombinant techniques in *E. coli*.

Two *E. coli* expression plasmids were constructed that expressed either the COOH-terminal 423 or 571 amino acid residues of the kinase in the Pinpoint  
30 fusion protein expression/purification system (Promega, Madison, Wisconsin). Briefly, DNA sequences encoding the COOH-terminal portion of the kinase

-26-

(nucleotides 1339 to 2630 or nucleotides 898 to 2630 of SEQ ID NO: 3) were fused in frame to the COOH-terminus of a 13 kD peptide derived from the transcarboxylase complex from propionibacterium shermanii. This region undergoes biotination in *E. coli*, and thus provides a means for monitoring expression and purification of the fusion proteins. Expression was driven from the tac promoter in pinpoint Xa3. Fusion protein expression was induced with 0.1mM IPTG and confirmed using streptavidin alkaline phosphatase in a pseudo-Western format as described by the manufacturer.

#### Example 4

Recombinant versions of MCCS1 may also expressed in yeast or in SF9 insect cells using a baculovirus expression system. The FRAP kinase has been expressed, purified and is enzymatically active after expression in the baculovirus system [Brown *et al.*, *supra*].

The coding region of MCCS1 is fused at the amino terminus to a heterologous peptide sequence, such as the FLAG tag MDYKDDDDK (SEQ ID NO: 20) or a six-histidine tag, and reconstructed into the appropriate vectors. Once expressed in insect cells, a monoclonal antibody that recognizes the FLAG tag (Eastman Kodak, Rochester, New York) is used to purify large quantities of the FLAG-PIK-related kinase fusion protein. Infected insect cells are incubated for 48 hours and lysed in lysis buffer (25mM 2-glycerolphosphate, 50mM sodium phosphate pH 7.2, 0.5% Triton-X 100, 2mM EDTA, 2mM EGTA, 25 mM sodium fluoride, 100μM sodium vanadate, 1mM PMSF, 1μg/ml leupeptin, 1μg/ml pepstatin, 1mM benzamidine, and 2mM DTT). Expressed FLAG fusion proteins are purified over a column containing anti-FLAG antibody M2 affinity resin (Eastman Kodak). The column is washed with 20 column volumes of lysis buffer, then 5 column volumes of 0.5M lithium chloride, 50mM Tris pH 7.6, 1mM DTT, and then eluted either with 0.1M glycine pH 3.0 followed by immediate neutralization or by competitive elution with the FLAG peptide. For six-histidine tagged proteins, Ni-NTA agarose (Qiagen) is used for protein purification.



-27-

Shortly after the filing of parent application U.S.S.N. 08/558,666, a gene identified as ATR was described by Antony M. Carr and co-workers (personal communications). ATR appears to encode the same or a closely related protein to MCCS1 based on a comparison of amino acid sequences between ATR and MCCS1. The DNA and deduced amino acid sequences of ATR are presented in SEQ ID NOs: 28 and 29, respectively. The sequence differences between ATR and MCCS1 $\beta$  are as follows. ATR includes an additional 98 amino acid residues at the N-terminus. At nucleotide position 1284 (SEQ ID NO: 32) there is a conservative base change from "A" in MCCS1 $\beta$  to "T" in ATR and at nucleotide position 4176, there is an additional conservative base change from "C" in MCCS1 $\beta$  to "T" in ATR.

The FLAG tag was fused at the amino-terminus of a truncated ATR molecule which lacked the first sixty-six ATR amino acids. The FLAG tag was added by PCR as follows. The oligos FLAG-ATR (5'-CGGGATCCGCCATGGACTACAAGGACGATGACAAGATGTTGCTTGATTTC-3'). And HFB24 (5'-CTTAAGCCGCATGAGCACACCGTC-3') were used in the following PCR reaction: 100ng of pcDNAATR (obtained from Antony M. Carr) as template; 1X PCR buffer (Perkin-Elmer Cetus); 1.5 mM MgCl<sub>2</sub>, 200 $\mu$ M each of dATP, dGTP, dCTP, and TTP, 10 ng/ $\mu$ l of each primer; 1U AmpliTaq (Perkin-Elmer Cetus). The reaction was denatured at 94°C for 4 minutes followed by 30 cycles of 94°C for 30 seconds, 60°C for 30 seconds and 72°C for 30 seconds. The resulting approximately 800 bp PCR product was digested with BamHI and NheI and was ligated to the 10kb fragment of the mammalian ATR expression plasmid, pcDNAATR digested with BamHI and BstXI along with the remainder of the ATR coding sequences contained on a 2.5 kb BstXI to NheI fragment. Sequence analysis confirmed the addition of the FLAG tag. The insert contained within this plasmid was then used to construct a baculovirus expression plasmid that would express the FLAG tagged ATR truncate. The 5' end of ATR contained on a BamHI to BstXI fragment and the 3' end of ATR contained on a BstXI to SalI fragment derived from pBTM ATR were ligated to the baculovirus expression vector, pFB (Gibco/BRL) that had been digested with BamHI and SalI. This plasmid was designated pFMBCCS $\beta$ FLAG.

-28-

The full coding region of ATR was fused at the amino terminus to the six histidine tag by PCR. Oligonucleotides MCCS6his (5'-CGGGATCCAGCATGCATCACCATCACCATCACATGGGGGAACATGGGC-3') and Frp1R (5'-CATGACCACTGGCCATTCCACACG-3') were used in a PCR reaction to add the six histidine tag to sequences encoding the amino-terminus of ATR. PCR conditions were as follows: 100 ng of PstA 12ATR (obtained from Antony M. Carr) was used as template; 1X PCR buffer (Perkin-Elmer Cetus); 1.5 mM MgCl<sub>2</sub>, 200 μM each of dATP, dGTP, dCTP, and TTP, 10 ng/μl of each primer; 1U AmpliTaq (Perkin-Elmer Cetus). The reaction was denatured at 94°C for 4 minutes followed by 25 cycles of 94°C for 30 seconds, 60°C for 30 seconds and 72°C for 30 seconds. The approximately 800 bp PCR product was digested with BamHI and MscI and ligated to two other fragments: a 10kb fragment from pcDNAATR digested with BamHI and BstXI and an approximately 3 kb MscI to BstXI fragment containing the remainder of the ATR coding sequence. The addition of the six histidine tag was verified by sequence analysis. The resulting plasmid encoding a six-histidine tagged full length ATR molecule was designated pcDNA6his ATR.

To construct a baculovirus expression plasmid that expressed the entire coding sequence of ATR, the 1.2 kb BamHI to AgeI fragment from pFBMCCSβFLAG was ligated to the BamHI to AgeI fragment from pcDNA6his ATR. The resulting plasmid, designated pFB/HisX6MCCS-1 plasmid was transformed into the E.coli strain, DH5α (Gibco/BRL) for screening of recombinants. This plasmid was purified by using the Promega "Wizard" mini-prep kit, then transformed into *E. coli* αSF9 cells (Invitrogen) using the Cellfectin protocol described by Gibco/BRL.

Forty eight hours after transfection, the SF9 cell pellet and baculovirus produced by the transfected cells were harvested. The virus was stored at 4°C in Grace's Complete media containing 10% FBS, Pennicillin-Streptomycin, and Gentamicin. This viral prep was used to make a high titer (P2) virus stock. The P2 virus stock was used to infect a 50 ml culture of SF9 cells. The cells were collected 48 hours after infection and centrifuged at low speed to pellet the cells without lysis. The cell pellet was stored at -20°C for 24 hours before lysis. The cells were lysed

-29-

in 5 ml of lysis buffer (50 mM Tris, pH 8.0; 500 mM NaCl; 1 % NP40; 100  $\mu$ M PMSF). Expression of ATR was confirmed by immunoblot using the polyclonal antibody anti-AgDH2 as a probe. The FBHisX6 ATR baculovirus produced an approximately 300 kDa protein that was immunoreactive with anti-AgDH2 antibodies and comigrated with a protein in a mouse testes cell extract.

The P2 virus stock was also used to infect a 2 liter culture of SF9 cells. The cells were collected 48 hours after infection, centrifuged at low speed to pellet the cells without lysis and stored at -20°C. A cell pellet from 150 mls of this culture was lysed in 7.5 ml of lysis buffer (50mM NaPO<sub>4</sub>, pH7.2; 0.5 % NP-40; 10mM imidazole, 25mM NaF, 100 $\mu$ M Na<sub>3</sub>VO<sub>4</sub>; 0.5mM AEBSF; 1  $\mu$ g/ml leupeptin; 1 $\mu$ g/ml pepstatin A) and incubated on ice for 15 minutes. The lysate was then centrifuged for 30 minutes at 10,000 x g. The supernatant was removed and any DNA in the lysate resulting from broken nuclei was sheared by aspirating through an 20 gauge needle. Particulate matter was then removed by filtering through a 0.8 micron filter followed by a 0.2 micron filter. This cleared lysate was adjusted to contain 5 mM  $\beta$ -mercaptoethanol and 0.4 M NaCl. A 1 ml Ni-NTA-agarose column (Qiagen) was equilibrated in Buffer A (0.4 M NaCl; 5 mM  $\beta$ -mercaptoethanol; 0.1 % Triton X-100; 50 mM NaPO<sub>4</sub> 10 mM imidazole; 25 mM NaF, 100  $\mu$ M Na<sub>3</sub>VO<sub>4</sub>; 0.5 mM AEBSF; 1  $\mu$ g/ml leupeptin; 1  $\mu$ g/ml pepstatin A) prior to loading the cleared lysate. The sample was loaded at a flow rate of 0.25 ml/minute, washed 5 ml of Buffer A and then eluted in 10 ml of a gradient of 50 to 500 mM imidazole in Buffer A. One half ml fractions were collected and was assayed for kinase activity as follows. Five  $\mu$ l of each fraction was incubated in kinase buffer, 10  $\mu$ Ci <sup>32</sup>P $\gamma$ ATP, 10  $\mu$ M ATP, and 5  $\mu$ g of substrate PHAS-1 (Stratagene) and incubated at 37°C for 20 minutes. The reaction was then spotted onto phosphocellulose spin columns and centrifuged at 2500x g, washed twice with 0.5 ml of 75 mM phosphoric acid and once with 0.5 ml absolute ethanol. The phosphocellulose disks were then transferred to scintillation vials and the counts per minutes incorporated into the PHAS-1 proteins were recorded. Fractions 4 through 9 were found to contain activity toward PHAS-1 and immunoblot analysis confirmed that ATR was also present in the same fractions.

-30-

MCCS1 encoding plasmid DNA was transformed into an *esr1-1* diploid yeast strain (*Mata leu2-1 his4-4 can1 ura3 cyh2 ade6 ade2 esr1-1/MAT a leu2-27 his4 trp1 met2 ade2 esr1-1*), and cells were grown to mid-log phase in either galactose or glucose containing medium. Cells were pelleted, washed and all steps performed at 4°C. Cell pastes were resuspended in buffer (20 mM Tris at pH 8.0, 300 mM NaCl, 10% glycerol, 0.1 mM PMSF, 0.25 mg/ml pepstatin, leupeptin, and aprotinin) and lysed in a French Press or using glass beads. Lysis was verified by microscopy following a low-speed (10K) spin and a high-speed spin (100K), and the supernatant was loaded onto a 1.5 ml Ni-NTA agarose (Qiagen, Inc., Chatsworth, CA) column prewashed in 1x buffer. The column was washed with six column volumes of buffer. The column was eluted stepwise with 8 ml of 10 mM, 50 mM, 100 mM, and 250 mM imidazole in buffer. Fractions were collected and Western analysis was performed using 15 µl of each elution peak. Kinase activity was measured as described above.

### Example 5

Polyclonal and monoclonal antibodies specific for MCCS1 were generated by standard techniques in the art.

Two different bacterial expression plasmids, pGEX1-MEC and pGEX3-MEC, were constructed for the recombinant production of portions of the MCCS1 polypeptide as fusions to the COOH-terminus of glutathione S-transferase (GST). Both plasmids were used for the generation of antigens AgDH-2 and AgDH-3, from pGEX1-MEC and pGEX3-MEC respectively for use in a standard immunization protocol. pGEX1-MEC contains an EcoRI fragment encoding amino acid residues 566 to 870 of SEQ ID NO: 4 fused to GST in the pGEX1 vector (Pharmacia Biotech, Milwaukee, Wisconsin); pGEX3-MEC contains an Eco RI fragment encoding amino acid residues 118 to 567 of SEQ ID NO: 4 fused to GST in the pGEX3 vector (Pharmacia Biotech). Induction of the pGEX tac promoter with 0.1mM IPTG led to high level expression of each fusion protein in an insoluble form (inclusion bodies). Following lysis of induced cultures with a French pressure cell, AgDH-2 and AgDH-3 extracts were centrifuged through a 35 % sucrose solution containing 0.1M NaCl,

-31-

0.01M Tris pH7.5, and 0.001M EDTA (STE). Pellets were then washed twice and resuspended in STE.

For the generation of polyclonal antisera in rabbits, AgDH-2 and AgDH-3 were further purified using preparative SDS polyacrylamide gel electrophoresis and electroelution of each antigen from gel slices. Primary immunization of female New Zealand White rabbits was with 200  $\mu$ g of each antigen mixed with complete Freund's adjuvant injected at multiple sites subcutaneously. Subsequent immunizations were with 100  $\mu$ g antigen mixed with incomplete Freund's adjuvant at approximately 21 day intervals, and test bleeds were taken after immunizations 3, 4 and 5. Western blot analysis of extracts of human testis tissue demonstrates antibody reactivity against an approximately 270 kD protein in immune but not preimmune antisera. In addition, the immune sera showed reactivity against the MCCS1 pinpoint fusion proteins described in Example 3, providing evidence of the generation of MCCS1-specific antibodies.

The MCCS1-specific antibodies were purified as follows. Inclusion body preparations of AgDH-2 and AgDH-3 were coupled to cyanogen bromide (CNBr)-activated Sepharose (Pharmacia, Alameda, CA). Two mg of antigen were solubilized in 1% SDS (4.5 ml final volume) and dialyzed overnight against Coupling Buffer (0.1M NaHCO<sub>3</sub>/0.1% SDS). 0.5 ml of 5M NaCl were added to each antigen preparation prior to incubation with the CNBr Sepharose. 0.4 gm of freeze-dried CNBr Sepharose (per antigen) were resuspended in 1 mM HCl and washed in a scintered glass funnel with 250 ml 1 mM HCl added in several aliquots over 15 minutes. The HCl-washed CNBr Sepharose was then removed to a 15 ml snap cap tube and washed twice with 5 ml of Coupling Buffer. Dialyzed antigen preps were added to the washed Sepharose and then incubated at room temperature for 1.5 hours on a slowly rotating wheel. The Sepharose was washed once with 5 ml of Coupling Buffer, once with 10 ml of 0.1M Tris pH8.0, and then incubated in 10 ml 0.1M Tris 8.0 for 2 hours at room temperature to block any remaining reactive groups on the resin. Coupling efficiency was 60-80% as judged by SDS-PAGE analysis. The antigen columns were then washed with 15 ml of 6M Guanidine HCl (to remove uncoupled antigen), 25 ml of Buffer A (50mM Tris pH 7.4), 15 ml of Buffer B (4.5M MgCl<sub>2</sub>/1mg/ml BSA/50mM Tris 7.4), and then 50 ml of Buffer A. Thirty ml

-32-

of rabbit serum from immunized animals (rabbit 4747 immunized with AgDH-3 and rabbit 4779 immunized with AgDH-2) were passed over the appropriate antigen column over the course of 3 hours. The columns were then washed with 20 ml of Buffer A, 40 ml of 1M Guanidine HCl, and then equilibrated with an additional 20 ml of Buffer A. Anti-AgDH-3 or Anti-AgDH-2 antibodies were then eluted off the antigen columns with 10 ml of Buffer B. One ml fractions were collected, IgG-containing fractions were pooled and dialyzed against 1 L of phosphate buffered saline (PBS) for 3 hours, and then overnight against 1 L of PBS containing 35% glycerol.

Antipeptide antibodies were generated against the human ATM protein by coupling a 15-amino-acid peptide (residues 1359-1373) to Keyhole Limpet Hemocyanin-using EDC as described by the manufacturer (Pierce), followed by injection of the coupled immunogen into rabbits. The antibodies were first precipitated from the serum (#6076) with an equal volume of saturated ammonium chloride followed by resuspension and dialysis against PBS. Affinity purification was carried out using a peptide column prepared by coupling the antigenic peptide to CNBr-activated Sepharose (Pharmacia) as described by the manufacturer. The antibodies were then bound to the peptide column and washed with 2 M KCl-PBS. Elution was carried out with 20 ml 5 M NaI (in 1 mM sodium thiosulfate), which was dialyzed immediately against PBS.

To generate monoclonal antibodies, female Balb/c mice were immunized with 50 ug AgDH-2 or AgDH-3. Additional mice were immunized with 25 to 50 ug AgDH-2 or AgDH-3 that had been combined with an equal molar ratio of mAb 61F3B, a monoclonal antibody with specific reactivity to GST. A third group of mice were immunized with SDS polyacrylamide gel slices containing AgDH-2 or AgDH-3. The immunogen for each group of mice was prepared in complete Freund's adjuvant, with subsequent boosts (25 ug antigen in incomplete Freund's) at about 21 day intervals. Cell lines producing monoclonal antibodies were isolated as follows. Briefly a single cell suspension was formed by grinding immunized mouse spleen in serum free RPMI 1640, supplemented with 2mM L-glutamine, 1mM sodium pyruvate, 100 units/ml penicillin, and 100 µg/ml streptomycin (RPMI) (Gibco, Canada). The cell suspension was filtered through

-33-

sterile 70-mesh Nitex cell strainer (Becton Dickinson, Parsippany, New Jersey), and washed twice by centrifuging at 200 g for 5 minutes and resuspending the pellet in 20 ml serum free RPMI. Thymocytes taken from three naive Balb/c mice were prepared in this manner.

5                   NS-1 myeloma cells kept in log phase in RPMI with 11 % fetal bovine serum (FBS) (Hyclone Laboratories, Inc., Logan, Utah) for three days prior to fusion, were centrifuged at 200 g for 5 minutes, and the pellet was washed twice as described in the foregoing paragraph. After washing, each cell suspension was brought to a final volume of 10 ml in serum free RPMI, and 10  $\mu$ l was diluted  
10               10:100. Twenty  $\mu$ l of each dilution was removed, mixed with 20  $\mu$ l 0.4% trypan blue stain in 0.85% saline (Gibco), loaded onto a hemacytometer and counted.

                  Two  $\times 10^8$  spleen cells were combined with  $4 \times 10^7$  NS-1 cells, centrifuged, and the supernatant was aspirated. The cell pellet was dislodged by tapping the tube and 2 ml of 37°C PEG 1500 (50% in 75mM Hepes, pH 8.0)  
15               (Boehringer Mannheim) was added with stirring over the course of 1 minute, followed by adding 14 ml of serum free RPMI over 7 minutes. An additional 16 ml RPMI was added and the cells were centrifuged at 200 g for 10 minutes. After discarding the supernatant, the pellet was resuspended in 200 ml RPMI containing  
20               15% FBS, 100  $\mu$ M sodium hypoxanthine, 0.4 $\mu$ M aminopterin, 16 $\mu$ M thymidine (HAT) (Gibco), 25 units/ml IL-6 (Mallinckrodt, Folcroft, Pennsylvania), and  $1.5 \times 10^6$  thymocytes/ml. The suspension was dispensed into ten 96-well flat bottom tissue culture plates at 200  $\mu$ l/well. Cells in plates were fed 3 to 4 times between fusing and screening by aspirating approximately half the medium from each well with an 18 G needle and replenishing plating medium described above except containing 10  
25               units/ml IL-6 and lacking thymocytes.

                  Fusions were screened when cell growth reached 60-80% confluency (day 7 to 9) by ELISA on AgDH2 versus AgDH3. Immulon 4 plates (Dynatech, Cambridge, MA) were coated at 4°C overnight with 100 ng/well protein in 30mM carbonate buffer, pH 9.6. Plates were blocked with 100  $\mu$ g/well 0.5% fish skin  
30               gelatin in PBS for one hour at 37°C, washed 3 times with PBS, 0.05% Tween 20 (PBST) and 50  $\mu$ l culture supernatant is added. After incubation at 37°C for 30 minutes, and washing as described above, 50  $\mu$ l of horseradish peroxidase conjugated

-34-

goat anti-mouse IgG(fc) (Jackson ImmunoResearch, West Grove, PA) diluted 1:10,000 in PBST was added. Plates were incubated as above, washed 4 times with PBST and 100  $\mu$ l substrate consisting of 100  $\mu$ g/ml of tetramethylbenzidine and 0.15  $\mu$ l/ml  $H_2O_2$  in 100mM sodium acetate, pH 5.5, was added. The color reaction was  
5 stopped in 5-10 minutes with the addition of 50  $\mu$ l of 15 %  $H_2SO_4$ .  $A_{490}$  was read on a plate reader.

Fifty three pools of hybridomas that were positive in an ELISA were screened for the ability to immunoblot or immunoprecipitate MCCA from a mouse testes cell lysate. Immunoblot analysis using the mouse testes extract is described in  
10 Example 6. Immunoprecipitations was performed as follows. A six percent SDS polyacrylamide gel was run and transferred to Immobilon-PVDF in 192 mM glycine, 25 mM Tris base, 0.1 % SDS, 20 % methanol, then blocked for 1 hour in 5 % powdered nonfat milk, 20 mM Tris pH 7.5, 100 mM NaCl 0.1 % Tween 20, and cut into the appropriate number of strips. The primary antibody (well supernatant) was  
15 diluted in the above block solution and incubated for one hour at room temperature, washed four times in block minus milk, incubated in goat anti-mouse IgG (H+L) HRP (BioRad #170-6516), washed again in block solution minus milk, transferred to NEN Renaissance ECL reagent and developed for 5 minutes.

Immunoprecipitation was performed as follows. Fifty  $\mu$ l of hybridoma supernatant was incubated for one hour on ice with 300  $\mu$ g of testes cell lysate prepared as described in Example 6. Thirty  $\mu$ l of a 50 % slurry of protein A agarose (Pierce, Rockford, IL), prebound to a rabbit anti-mouse bridging antibody (5  $\mu$ g/reaction) (Pierce) was added and incubated at 4°C with rocking. The immune complexes were washed three times in lysis buffer and the antigen/antibody complex  
25 eluted by boiling in SDS sample buffer (2 % SDS, 20 mM Tris pH 6.8, 20 % glycerol, 0.001 % bromphenol blue). The resulting supernatant was separated on a 6 % SDS polyacrylamide gel and transferred to Immobilon-PVDF (Millipore) and an immunoblot was performed using affinity purified rabbit anti-Ag DH2 polyclonal antiserum. Four hybridomas were cloned and characterized in immunoblots,  
30 immunoprecipitations and in immunoprecipitation/kinase assays as described in Example 6. The four hybridoma cell lines were designated 224B, 224C (ATCC HB



-35-

12233), 224F (ATCC HB 12234) and 224G. All four monoclonal antibodies recognized MCCS1 by immunoblot and immunoprecipitation.

### Example 6

5 MCCS1 associated protein kinase activity was immunoprecipitated using the MCCS1-specific polyclonal antibodies described in Example 5.

Extracts were made from fresh testes tissue isolated from Balb/c mice. Minced testes were homogenized on ice with 10-15 strokes of a tight fitting dounce homogenizer in Lysis Buffer (50 mM NaPO<sub>4</sub>, pH 7.2; 0.5 % TritonX-100; 2 mM EDTA; 2 mM EGTA; 25 mM NaF; 25 mM 2-glycerophosphate; 1 mM  
10 phenylmethylsulfonyl fluoride [PMSF]; 1 µg/ml leupeptin; 1 µg/ml pepstatin A; 2 mM DTT) and incubated on ice for 30 minutes. The lysate was centrifuged at 13,000xg rpm for 10 minutes at 4° C in a TL-100 table-top ultracentrifuge (Beckman) to remove unbroken cells and other insoluble material. Aliquots of cell lysate were snap frozen in liquid N<sub>2</sub> and stored at -70°C. Five hundred ug of testes extract was  
15 incubated with either 5 ug of affinity purified anti-AgDH-2 polyclonal antibody or 5 ug purified rabbit IgG (Zymed, So. San Francisco, CA) in 1 ml of Lysis buffer for one hour on ice in microcentrifuge tubes. Thirty µl of protein A sepharose beads (Repligen, Cambridge, MA) (washed in Lysis buffer) were added to the extracts, and then incubated for an additional 30 minutes at 4° C on a rocking platform. The  
20 immune complex/Protein A sepharose beads were washed four times with 1 ml of Lysis buffer, one time with 1 ml Kinase Buffer (25 mM Hepes pH 7.7; 50 mM KCl; 10 mM MgCl<sub>2</sub>; 0.1 % NP-40; 2 % glycerol; 1 mM DTT), and then incubated in 20 µl Kinase Buffer with 10 µCi ATP [50 Ci/mmol] for 20 minutes at 37°C. The kinase reactions were stopped with 20 µl 2X SDS sample buffer and heated to 100°  
25 C prior to separation on 6% polyacrylamide gels. Gels were fixed in 20% methanol/7% Acetic acid, and then dried onto Whatman 3MM paper prior to autoradiography. While little or no phosphorylation was evident in control immunoprecipitations, immunoprecipitations using anti-AgDH-2 antibody contained two major phosphorylated bands at approximately 300 kD and approximately 180 kD.  
30 In addition, there were several minor phosphorylation products, including one which comigrated with the MCCS1 protein itself as demonstrated by Western blot analysis

-36-

(see Example 8 for Western blot description.) Phosphoaminoacid analysis of the approximately 300 kD protein identified the presence of phosphoserine residues. Addition of 5 ug of AgDH-2 (but not AgDH-3) dramatically reduced or eliminated the MCCS1-associated kinase activity found in the immunoprecipitates.

5

### Example 7

The expression pattern of MCCS1 in various human tissues was examined by Northern blot hybridization.

Nylon membranes containing 2  $\mu$ g of size-fractionated polyA+ RNA from a variety of human tissue sources were obtained from Clontech Laboratories, Inc., and the hybridization protocol supplied by the manufacturer was followed precisely, except that the final wash was performed at 55° C, rather than 50° C, to minimize the possibility of cross-hybridization to related sequences. The <sup>32</sup>P-labelled DNA hybridization probe used was generated by PCR. A DNA encoding the COOH-terminal 30% of MCCS1 $\alpha$  was used as a template to amplify a 1.3 kb fragment in the presence of <sup>32</sup>P-dCTP using primers 279-3 5'TGGATGATGACAGCTGTGTC 3' (SEQ ID NO: 21) and 279-6 5'TGTAGTCGCTGCTCAATGTC3' (SEQ ID NO: 22).

Results of the Northern blots show that MCCS1 is expressed as an approximately 9 kb mRNA in a wide variety of human tissues. Testis tissue contains the highest level of MCCS1 mRNA, though the transcript is also expressed in small intestine, ovary, prostate, thymus, spleen, heart, peripheral blood lymphocytes, colon, brain, placenta, skeletal muscle, kidney and pancreas.

Expression of MCCS1 mRNA in human cancer cell lines was also examined using a human cancer cell line RNA blot obtained from Clontech. The RNA blot contained RNA from the cell lines HL-60 (promyelocytic leukemia), HeLa (cervical carcinoma), K-562 (chronic myelogenous leukemia), MOLT-4 (lymphoblastic leukemia), Raji (Burkitt's lymphoma), SW480 (colorectal adenocarcinoma), A549 (lung carcinoma), and G361 (melanoma). Northern blot analysis was performed as directed by the manufacturer with hybridization being carried out at 65°C using a 2.0kb KpnI-SalI fragment of the MCCS1 partial clone HFB2. Expression was observed in the HL-60, HeLa, K-562, Raji, SW480, and G361 cell lines with the highest level of expression occurring in the G361 cell line.

-37-

Detectable but low levels of expression were observed in the MOLT-4 and A549 cell lines.

### Example 8

5 The expression of MCCS1 mRNA and protein in normal and irradiated mouse testes and in mouse embryos was examined by *in situ* hybridization, immunostaining and/or immunoblotting.

#### In situ Hybridization

Normal and irradiated mouse testes were harvested from male Balb/c mice. The tissues were sectioned at 6 $\mu$ m thickness, picked up on Superfrost Plus<sup>®</sup> (VWR Scientific) slides and allowed to air-dry at room temperature overnight. Sections were  
10 stored at -70 $^{\circ}$  C if not immediately used. The tissue sections were fixed in 4% paraformaldehyde (Sigma) in PBS for 20 minutes at 4 $^{\circ}$  C, dehydrated (70%, 95%, 100% ethanol) for 1 minute at 4 $^{\circ}$  C in each grade, then allowed to air dry for 30 minutes at room temperature. The slides were acetylated in a solution of 0.25% (v/v)  
15 acetic anhydride (Sigma)/0.1M triethanolamine pH 8.0 for 10 minutes at room temperature with stirring, rinsed in 0.2X SSC for 10 minutes at room temperature with stirring, and dehydrated and air dried as described above. The tissues were hybridized *in situ* with digoxigenin-labeled single-stranded mRNA generated from murine MCCS1 DNA by *in vitro* RNA transcription incorporating digoxigen-UTP  
20 (Boehringer Mannheim). The labeled riboprobes (see sequence in SEQ ID NO: 27) (1 $\mu$ g/section) and diethylpyrocarbonate (depc)-treated water were added to hybridization buffer with a final concentration of 50% formamide, 0.3 M NaCl, 20 mM Tris pH 7.5, 10% dextran sulfate, 1X Denhardt's solution, 100 mM dithiothreitol (DTT) and 5 mM EDTA, and 20  $\mu$ l of the solution was applied to each  
25 section and covered with a sterile, RNase-free 22 x 22 cover slip. The mRNA in both the section and the probe solution was denatured by heating the slides to 85 $^{\circ}$  C for 10 minutes in an oven. Hybridization was carried out overnight (12-16 hours) at 50 $^{\circ}$  C.

After hybridization, sections were washed for 1 hour at room  
30 temperature in 4X SSC/10 mM DTT, then for 30 minutes at 50 $^{\circ}$  C in 50% formamide/2X SSC/10 mM DTT, 30 minutes at 37 $^{\circ}$  C in a solution of 500 mM

-38-

NaCl, 10 mM Tris-HCl, 1 mM EDTA, pH 7.5 (NTE buffer), 30 minutes at 37° C in a bath of 10 µg/mL RNase A (Boehringer Mannheim) in NTE buffer, 15 minutes at 37° C in NTE buffer, 15 minutes at room temperature in 2X SSC, 15 minutes at room temperature in 0.1X SSC, and 2 minutes at room temperature in 100 mM Tris-HCl, 150 mM NaCl, pH 7.5 (Buffer 1). To detect the labeled riboprobes, the sections were blocked for 30 minutes at room temperature in a solution of 5% normal sheep serum (Harlan Bioproducts for Science, Indianapolis, IN) and 0.3% Triton X-100 (Sigma) in Buffer 1 with gentle stirring, after which 150 µl/section of sheep αDigoxigenin-gold conjugate (Goldmark Biologicals, Philipburg, Pa) was applied to the tissues and incubated for 2 hours at room temperature. The slides were then washed three times for 5 minutes in Buffer 1, five times for 3 minutes in sterile deionized water, the excess liquid blotted off the slide and 2 drops each of silver enhancing and initiating solution (Goldmark Biologicals) applied to each section. The chemical reaction was allowed to proceed for 23 minutes at room temperature, then the sections were rinsed thoroughly in sterile deionized water, counterstained in Nuclear Fast Red (Vector), rinsed again in sterile deionized water, air dried overnight at room temperature and mounted with Cytoseal 60 (VWR).

In both normal and irradiated mouse testes signal was observed in the cytoplasm of spermatogonia and spermatocytes. The expression level in irradiated testis was not increased over that seen in normal testis.

#### Immunostaining

Testis tissue from normal male Balb/c mice was sectioned at 6 µm thickness, picked up on Superfrost Plus® (VWR Scientific) slides and allowed to air-dry at room temperature overnight. Sections were stored at -70° C if not immediately used. The sections were fixed in cold (4° C) acetone for 10 minutes at room temperature; once the slides were removed from the acetone the reagent was allowed to evaporate from the sections. Each tissue section was blocked with 150 µl of a solution of 30% normal rat serum (Harlan Bioproducts), 5% normal goat serum (Vector Laboratories) and 1% bovine serum albumin (BSA) (Sigma) in 1X TBS for 30 minutes at room temperature. After blocking, the solution was gently blotted from the sections and anti-AgDH-3 and anti-AgDH-2 polyclonal antibodies and preimmune sera from the same rabbits were diluted 1:50 and 1:100 in the blocking solution and

-39-

100  $\mu$ l applied to each tissue section and incubated for 30 minutes at 37° C. The antibody solution was blotted gently from the sections and unbound antibody removed from the sections by washing the slides 3 times for 5 minutes each in 1X TBS. The excess TBS was blotted from the slide and 100  $\mu$ l of the biotinylated goat anti-rabbit antibody contained in the Elite Rabbit IgG Vectastain ABC kit (Vector), prepared according to the product insert, were applied to each section and incubated for 15 minutes at 37° C. After incubation, the slides were washed 2 times in 1X TBS for 5 minutes in each wash. Next, 100  $\mu$ l of streptavidin-gold conjugate (Goldmark Biologicals) diluted 1:100 in a solution containing 5% normal rat serum and 1% BSA was applied to each section and incubated for 1 hour at room temperature. The slides were then washed 3 times in 1X TBS for 5 minutes each wash, and 100  $\mu$ l of 1% glutaraldehyde (Sigma) in TBS buffer was applied to the slides for 5 minutes at room temperature. The slides were then washed 3 times for 5 minutes each in TBS, then 4 times in sterile deionized water for 3 minutes each. The excess liquid was blotted from each slide and 2 drops each of silver enhancing and initiating solution (Goldmark Biologicals) were applied to each section. The chemical reaction was allowed to proceed for 13 minutes at room temperature, then the sections were rinsed thoroughly in sterile deionized water, counterstained in Nuclear Fast Red (Vector), rinsed again in sterile deionized water, air dried overnight at room temperature and mounted with Cytoseal 60 (VWR).

Signal was detected in the spermatogonia and primary spermatocytes with both of the polyclonal antibodies, but not with the preimmune sera from the same animals.

#### Immunoblotting

Freshly obtained mouse testicles were minced with razor blades in cold PBS, and a cell suspension was generated using a loose fitting dounce homogenizer. This cell suspension was then boiled with an equal volume of 2X SDS sample buffer. Fifty  $\mu$ g aliquots of each extract were separated on 6% polyacrylamide gels, transferred onto Immobilon membranes (Millipore, Bedford, MA) and analyzed for anti-MCCS1-reactivity using the affinity purified antibodies in Example 5, and HRP-conjugated goat anti-rabbit secondary antibody and the Renaissance Enhanced Chemiluminescence kit (Dupont/NEN, Boston, MA). Extracts prepared from fresh

-40-

mouse testis contain a high molecular weight species (about 294 kD) that was recognized by both affinity-purified antiserum. No reactivity against this protein was seen with either of the preimmune sera. Importantly, the signal obtained from each affinity purified sera was specifically blocked after pre-incubation of the antibody with the corresponding immunogen.

In summary, high levels of M CCS1 mRNA and protein are detected in mouse testis in the spermatogonia and primary spermatocytes, cells that are in the early stages of meiosis. This suggests that M CCS1 plays an important role in meiotic cell division. Meiosis is a specialized form of cell division that produces germ cells in higher eukaryotes. There are two major characteristics of meiosis that distinguish it from mitosis. Whereas mitotic cell division results in genetically identical cells containing two of each chromosome, meiotic cell division results in cells containing one of each chromosome. Early in meiosis, during the "reduction division" process, sister chromatids pair and undergo reciprocal recombination at some regions. During this process, these cells are exposed to DNA strand breaks. It is likely that the cellular response to the DNA strand breaks during meiosis is similar to the cellular response found in non-germ cells in response to IR-induced DNA damage. This interpretation is further substantiated by studies that demonstrate the MEC1 is upregulated 10 to 20 fold during sporulation, indicating an important role for M CCS1 during meiosis in addition to its role in DNA repair.

#### Example 9

In order to identify the cells within the developing mouse testis that express M CCS1, Western blot analysis of M CCS1 expression within populations of meiotic cells was performed. Extracts of purified pachytene spermatocytes, round spermatids, condensing spermatids, and epididymal sperm cells were examined for M CCS1 expression as described above in Example 8.

Pachytene spermatocytes, round, and condensing spermatids were prepared from decapsulated testes of adult mice by sequential dissociation with collagenase and trypsin-DNase 1. The cells were separated into discrete populations by sedimentation velocity at unit gravity in 2-4% BSA gradients in Enriched Krebs Ringer Bicarbonate Medium (EKRB). The pachytene spermatocyte and round

-41-

spermatid populations were each at least 85% pure, while the condensing spermatid population was about 40-50% pure (contaminated primarily with enucleated residual bodies and some round spermatids). Sperm were obtained from the cauda epididymides. Purified populations of spermatogenic cells were dissolved directly in SDS-sample buffer containing 40 mM DTT, heated to 100° C for 5 minutes, and the amount of protein in each sample determined by the Amido-Black procedure.

The highest levels of MCCS1 protein were found in pachytene spermatocytes, with the level dropping significantly in round spermatids. MCCS1 protein levels were barely detectable lower in the condensing spermatid population, and this may reflect the presence of round spermatids in the preparation (see above). No MCCS1 protein was detected in epididymal sperm. The Western analysis thus corroborates the immunocytochemical data, and suggests a role for MCCS1 in meiotic cells.

#### Example 10

Substrates of MCCS1 and proteins that interact with MCCS1 (for example, members of the cell cycle checkpoint pathway and proteins that localize MCCS1 in cells) may be identified by various assays.

##### A. Identification of Substrates

Substrates of MCCS1 may be identified by incorporating test compounds in assays for kinase activity. MCCS1 kinase is resuspended in 20  $\mu$ l kinase buffer (25mM Hepes pH7.4, 25mM KCl, 10mM MgCl<sub>2</sub>, 1mM DTT, 2% glycerol, 0.1% NP40, 0.5mM ATP, 10 uCi gamma <sup>32</sup>P-ATP) and incubated for 30 minutes, either in the presence or absence of 4  $\mu$ g test compound (e.g., casein, histone H1, or appropriate substrate peptide). Reactions are separated on 12% PAGE gels and dried onto Whatman paper prior to autoradiography. Moles of phosphate transferred by the kinase to the test compound are measured by autoradiography or scintillation counting. Transfer of phosphate indicates that the test compound is a substrate of the kinase.

The protein PHAS-1 has been identified as an *in vitro* substrate of ATR (Example 4). PHAS-1 is a heat and acid-stable protein that phosphorylated at several sites *in vivo* in response to insulin and growth factors. PHAS-1 binds to the mRNA

-42-

cap binding factor, EIF-4E, and prevents translation of capped mRNAs. Phosphorylation of PHAS-1 at a specific serine residue results in dissociation of PHAS-1 from EIF-4E and thus releasing the inhibition of translation of capped mRNAs. This mechanism allows for a rapid synthesis of protein in response to a particular stimulus. PHAS-1 may be phosphorylated by several protein kinases *in vivo* including a protein kinase that is sensitive to rapamycin. Since the rapamycin-sensitive protein kinase, FRAP, is related to ATR, it would be reasonable to assume that there might be an overlap in substrate specificity between FRAP and ATR and that PHAS-1 is a substrate for both of these protein kinases *in vitro*. To test this hypothesis, ATR that was immunoprecipitated from a mouse testes cell extract or His-tagged ATR purified from baculovirus-infected SF9 cells (Example 4) was incubated with 10  $\mu$ g PHAS-1 (Stratagene) in kinase buffer (25 mM Hepes pH 7.4, 25 mM KCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 0.1% NP-40), 10  $\mu$ M ATP and 10  $\mu$ Ci<sub>32</sub>P $\gamma$ ATP for 20 minutes at 37°C. Since phosphorylated PHAS-1 was known to bind to phosphocellulose paper, the reaction was spotted onto phosphocellulose spin columns and centrifuged at 2500 x g, washed twice with 0.5 ml of 75 mM phosphoric acid and once with 0.5 ml absolute ethanol. The phosphocellulose disks were then transferred to scintillation vials and the counts per minutes incorporated into the PHAS-1 proteins were recorded. ATR readily phosphorylated PHAS-1 whereas negative controls showed little or no PHAS-1 phosphorylation. To map which residue is phosphorylated, the following peptides representing PHAS-1 sequences containing serine and threonine residues were synthesized.

Peptide PH-1

MSGGSSCQTPSRAIPATRR (SEQ ID NO: 36)

Peptide PH-2

GDYSTTPGGTLFSTTPGGTRR (SEQ ID NO: 37)

Peptide PH-3

ECRNSPVTKTTRR (SEQ ID NO: 38)

Peptide PH-4

GVTSPSSDEPRR (SEQ ID NO: 39)

Peptide PH-5

MEASQSHLRR (SEQ ID NO: 40)



-43-

Peptide PH-6

RRNSPEDKRAGG (SEQ ID NO: 41)

Peptide PH-7

GEESQFEMDIRR (SEQ ID NO: 42)

5        These peptides are tested in the same kinase reaction to determine which peptide(s) is (are) phosphorylated by ATR. The peptide(s) are then used as substrate for ATR or MCCS1 in assays such as described in Example 11 to identify modulators.

         The same kinase reaction was also used to determine if proteins such as histone H1 (Upstate Biotechnology, Inc., Waltham, NY) and myelin basic protein  
10        (Gibco BRL, Gaithersburg, MD) which are known to be substrates of other protein kinases are substrates of MCCS1 and ATR. No phosphorylation of histone H1 or myelin basic protein was observed under the conditions of the assay. Moreover, a peptide from p53 known to be a substrate of DNA-PK was also not phosphorylated in the assay.

15        B. Identification of Interacting Proteins

         Interacting proteins may be identified by the following assays.

         A first assay contemplated by the invention is a two-hybrid screen. The two-hybrid system was developed in yeast [Chien *et al.*, *Proc. Natl. Acad. Sci. USA*, 88: 9578-9582 (1991)] and is based on functional *in vivo* reconstitution of a  
20        transcription factor which activates a reporter gene. Specifically, a polynucleotide encoding a protein that interacts with MCCS1 is isolated by: transforming or transfecting appropriate host cells with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA binding domain and an activating domain; expressing in the host cells a first hybrid  
25        DNA sequence encoding a first fusion of part or all of MCCS1 and either the DNA binding domain or the activating domain of the transcription factor; expressing in the host cells a library of second hybrid DNA sequences encoding second fusions of part or all of putative MCCS1 binding proteins and the DNA binding domain or activating domain of the transcription factor which is not incorporated in the first fusion;  
30        detecting binding of an MCCS1 interacting protein to MCCS1 in a particular host cell by detecting the production of reporter gene product in the host cell; and isolating second hybrid DNA sequences encoding the interacting protein from the particular

-44-

host cell. Presently preferred for use in the assay are a *lexA* promoter to drive expression of the reporter gene, the *lacZ* reporter gene, a transcription factor comprising the *lexA* DNA binding domain and the GAL4 transactivation domain, and yeast host cells.

5 Other assays for identifying proteins that interact with M CCS1 may involve immobilizing M CCS1 or a test protein, detectably labelling the nonimmobilized binding partner, incubating the binding partners together and determining the amount of label bound. Bound label indicates that the test protein interacts with M CCS1.

10 Another type of assay for identifying M CCS1 interacting proteins involves immobilizing M CCS1 or a fragment thereof on a solid support coated (or impregnated with) a fluorescent agent, labelling a test protein with a compound capable of exciting the fluorescent agent, contacting the immobilized M CCS1 with the labelled test protein, detecting light emission by the fluorescent agent, and identifying  
15 interacting proteins as test proteins which result in the emission of light by the florescent agent. Alternatively, the putative interacting protein may be immobilized and M CCS1 may be labelled in the assay.

### Example 11

Modulators of M CCS1 include M CCS1 variants and other molecules.  
20 The modulators may affect M CCS1 kinase activity, its localization in the cell, and/or its interaction with members of the cell cycle checkpoint pathway. Presently preferred regions of M CCS1 which are targets for mutation or the development of selective modulators include the following four regions: the M CCS1 $\alpha$  amino terminal effector domain (amino acids 1 to 1081 of SEQ ID NO: 31), the M CCS1 $\beta$  amino  
25 terminal effector domain (amino acids 1 to 1150 of SEQ ID NO: 33), the M CCS1 $\alpha$  *rad3*+ domain (amino acids 1082 to 2082 of SEQ ID NO: 31), the M CCS1 $\beta$  *rad3*+ domain (amino acids 1151 to 2151 of SEQ ID NO: 33), the M CCS1 $\alpha$  PIK domain (amino acids 2083 to 2410 of SEQ ID NO: 31), and the M CCS1 $\beta$  PIK domain (amino acids 2152 to 2480 of SEQ ID NO: 33).

30 M CCS1 variants having mutations in the kinase domain may be useful as a radiosensitizing agents. Mutations specifically contemplated by the invention are,

replacement of the MCCS1 $\alpha$  aspartic acid at amino acid 2241, the asparagine at 2246, and the aspartic acid at 2260 of SEQ ID NO: 31 with alanine or methionine, and the corresponding mutations in MCCS1 $\beta$ . Analogous mutations in the *rad3+* gene resulted in yeast hypersensitive to radiation. In addition, mutations in the kinase domain of ATM are found in patients with AT, a disease that causes radiation sensitivity.

Furthermore, combinatorial libraries, peptide and peptide mimetics, defined chemical entities, oligonucleotides, and natural product libraries may be screened for activity as modulators in assays such as those described below.

For example, an assay for identifying modulators of MCCS1 kinase activity involves incubating an MCCS1 kinase preparation in kinase buffer with gamma-<sup>32</sup>P-ATP and an exogenous kinase substrate, both in the presence and absence of a test compound, and measuring the moles of phosphate transferred to the substrate. For example, 2  $\mu$ l of the 50 mM imidazole elution pool is added to kinase buffer. (See Example 6.) The reactions are incubated at 37°C for 20 min and samples are analyzed by SDS-PAGE prior to autoradiography or Western analysis. An increase in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the test compound is an activator of said MCCS1 kinase. Conversely, a decrease in the moles of phosphate transferred to the substrate in presence of the test compound compared to the moles of phosphate transferred to the substrate in the absence of the test compound indicates that the modulator is an inhibitor of said MCCS1 kinase.

Moreover, assays for identifying compounds that modulate interaction of MCCS1 with other proteins may involve: transforming or transfecting appropriate host cells with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA-binding domain and an activating domain; expressing in the host cells a first hybrid DNA sequence encoding a first fusion of part or all of MCCS1 and the DNA binding domain or the activating domain of the transcription factor; expressing in the host cells a second hybrid DNA sequence encoding part or all of a protein that interacts with MCCS1 and the DNA binding domain or activating domain of the transcription factor which is not

-46-

incorporated in the first fusion; evaluating the effect of a test compound on the interaction between MCCS1 and the interacting protein by detecting binding of the interacting protein to MCCS1 in a particular host cell by measuring the production of reporter gene product in the host cell in the presence or absence of the test compound; and identifying modulating compounds as those test compounds altering production of the reported gene product in comparison to production of the reporter gene product in the absence of the modulating compound. Presently preferred for use in the assay are a *lexA* promoter to drive expression of the reporter gene, the *lacZ* reporter gene, a transcription factor comprising the *lexA* DNA binding domain and the GAL4 transactivation domain, and yeast host cells.

Another type of assay for identifying compounds that modulate the interaction between MCCS1 and an interacting protein involves immobilizing MCCS1 or a natural MCCS1 interacting protein, detectably labelling the nonimmobilized binding partner, incubating the binding partners together and determining the effect of a test compound on the amount of label bound wherein a reduction in the label bound in the presence of the test compound compared to the amount of label bound in the absence of the test compound indicates that the test agent is an inhibitor of MCCS1 interaction with protein. Conversely, an increase in the bound in the presence of the test compound compared to the amount label bound in the absence of the compound indicates that the putative modulator is an activator of MCCS1 interaction with the protein.

Yet another method contemplated by the invention for identifying compounds that modulate the binding between MCCS1 and an interacting protein involves immobilizing MCCS1 or a fragment thereof on a solid support coated (or impregnated with) a fluorescent agent, labelling the interacting protein with a compound capable of exciting the fluorescent agent, contacting the immobilized MCCS1 with the labelled interacting protein in the presence and absence of a test compound, detecting light emission by the fluorescent agent, and identifying modulating compounds as those test compounds that affect the emission of light by the fluorescent agent in comparison to the emission of light by the fluorescent agent in the absence of the test compound. Alternatively, the MCCS1 interacting protein may be immobilized and MCCS1 may be labelled in the assay.

### Example 12

Cell based complementation assays for identifying modulators of MCCC1 or ATM are described below.

5 In one type of assay, host cells (for example, *esr1-1* yeast cells) are transformed with MCCC1-encoding DNA as is described in Example 4. The *esr1-1* yeast strain is normally sensitive to treatment with ultraviolet (UV) light, but *esr1-1* yeast cells expressing MCCC1 or ATR are no longer sensitive to treatment with UV light. The transformed yeast cells are exposed to test compounds and the effect of the test compounds on UV sensitivity of the transformed host cell is determined. Test  
10 compounds that are inhibitors of MCCC1 or ATR activity restore UV sensitivity to the MCCC1 transformed *esr1-1* cells. Alternatively, *esr1-1 tel1* double mutant yeast cells are used as host cells instead of *esr1-1* yeast cells. The TEL1 gene is homologous to ATM and the TEL1 mutation is described in Morrow, *et al.*, *Cell*, 82:831-840 (1995). The invention also specifically contemplates that the *esr1-1* or  
15 *esr1-1 tel1* double mutant yeast host cells may be transformed with ATM-encoding DNA (SEQ ID NO: 34).

In an alternative embodiment, the assays include clastogenic agents or events instead of treatment with UV light (*e.g.*, IR, hydroxyurea, or DNA damaging agents). Appropriate host cells for use in such embodiments would be those that are  
20 sensitive to the alternative clastogenic agents or events.

Another type of complementation assay involves the use of mammalian host cells such as cell lines derived from cells of AT patients. As described above for yeast cells, the mammalian cells are transfected with DNA encoding MCCC1, ATR, or ATM and then exposed to test compounds. Test compounds that are  
25 inhibitors of MCCC1, ATR, or ATM activity will restore the phenotype of the untransformed host cell (*e.g.*, sensitivity to IR).

The above assays can be used to identify compounds that inhibit activity of MCCC1, ATR, and ATM or compounds that inhibit activity of only one of the enzymes.

30 In an alternative type of assay, the yeast or mammalian host cells are transformed with DNA encoding chimeric polypeptides including various combinations of MCCC1 and ATM domains. MCCC1 and ATM show structural

-48-

similarities, and chimeric polypeptides which comprise portions of MCCC1 and ATM are useful in elucidating active sites and binding domains of both MCCC1 and ATM. Polynucleotides encoding the chimeras can be prepared by standard molecular biology techniques known to the skilled worker and as exemplified herein. The chimeric polypeptides are expressed in host cells and modulators of the chimeras can be identified by the assays disclosed herein.

### Example 13

MCCC1 and ATM are both involved in meiosis I checkpoints. Since MCCC1 is demonstrated herein to have kinase activity, assays were performed to determine if ATM possessed kinase activity. To determine the kinase activity of ATM, ATM was immunoprecipitated from MRC-5 fibroblasts (ATCC #171-CCL) with polyclonal antisera, 6076. MRC-5 cells are human lung embryonal diploid fibroblasts. MRC-5 cells were obtained from the ATCC at passage 19 and maintained in Minimal Essential Medium supplemented with 10% fetal bovine serum, 100 units/ml penicillin, 100 mg/ml streptomycin, and 100 mM MEM non-essential amino acids. Media and media supplements were obtained through Gibco Life Technologies. Cell lines were maintained in a water-saturated 37°C incubator with 5% C.

MRC-5 cell extracts were prepared by lysis of a 10cm plate of log-phase cells in 0.5 ml of Lysis Buffer I (50 mM NaPO<sub>4</sub>, pH 7.2; 0.5% TritonX-100; 2 mM EDTA; 2 mM EGTA; 25 mM NaF; 25 mM 2-glycerophosphate; 1 mM phenylmethylsulfonyl fluoride [PMSF]; 1 µg/ml leupeptin 1 µg/ml pepstatin A; 2 mM DTT) on ice. Cells were scraped from plates using a rubber spatula then sonicated in a cup horn sonicator (Sonifier 250, Branson Ultrasonics Corp., Danbury, CT) at 100% output for 90 seconds. Lysates were then clarified in a 4°C microfuge for 2 minutes. Preclearing was done by adding 10 µg purified rabbit IgG (Zymed) and 30 µl Protein A Agarose slurry (Pierce) followed by incubation at 4°C for 60 minutes while rocking. To the precleared lysates, 10 µg of affinity purified 6076 antisera (or 10 µg 6076 pre-blocked with 0.04 mg P45 peptide for 30 min.) was added and incubated on ice for 60 minutes. Immunoprecipitates were collected by addition of

30  $\mu$ l Protein A agarose slurry and incubated with rocking at 4°C for 30 minutes followed by four washes in Lysis Buffer I.

Kinase reactions were carried out by washing the immunoprecipitations once with kinase buffer (25  $\mu$ M Hepes pH 7.7; 50 mM KCl; 10 mM MgCl<sub>2</sub>; 0.1 % NP-40; 2 % glycerol; 1 mM DTT), followed by incubation in 20  $\mu$ l of Kinase Buffer containing 10  $\mu$ M ATP + 10  $\mu$ Ci  $\gamma$  <sup>32</sup>P-ATP [50 Ci/mmol] for 20 minutes at 37°C. Reactions were stopped by the addition of 20  $\mu$ l 2X SDS sample buffer and boiled for 5 minutes prior to separation on 6% SDS polyacrylamide gels. The gels were dried and exposed to x-ray film (Kodak, XAR-5) at -80°C overnight.

10 10 cm plates of log-phase MRC-5 cells were washed once with PBS then incubated in Dulbecco's Modified Eagle Medium (minus methionine) containing 2% dialyzed fetal bovine serum for 30 minutes. Cells were labeled by adding 200  $\mu$ Ci <sup>35</sup>S-methionine (1000 Ci/mmol TRAN<sup>35</sup>S-LABEL, ICN Radiochemicals) for 2 hours. Labeled cells were then washed once with PBS and frozen at -80°C prior to immunoprecipitation.

The incubation of the immunoprecipitated complexes in kinase buffer produced a phosphorylated product with a molecular weight of approximately 350,000 that co-migrated with ATM in polyacrylamide gels.

20 Similar results were obtained for ATR immune complexes immunoprecipitated with anti-AgDH-2 (MCCS1) polyclonal antisera of Example 5. ATR and ATM thus appear to be able to self-phosphorylate or associate with a protein kinase.

25 To determine the role of ATR and ATM in meiosis, immunostaining techniques on surface spreads of mouse spermatocytes were utilized to localize ATR and ATM to meiotic chromosomes. Antibodies recognizing ATR and ATM were utilized with mouse antibodies against Cor1. Cor1 is a component of axial/lateral elements of synapsing chromosomes [Dobson *et al.*, *J. Cell Sci.*, 107:2749-2760 (1994)]. Cor1 chromosomal staining appears when the axial elements begin to form between the sister chromatids of each homolog in leptotema of meiotic prophase, prior to the initiation of synapsis. As homologous bivalents synapse, the axial elements from the two homologs align and a central element forms between them, completing the structure called the synaptonemal complex (SC).

-50-

When short stretches of Cor1 begin to appear prior to any evidence of synapsis, neither ATR nor ATM is detectable. As homologs start synapsis, both proteins were seen at pairing forks; however, the location and behavior of the two proteins differed markedly. In normal zygotene nuclei, the stage during which homologs synapse, ATR was present in small amounts and transiently at discrete foci along the asynapsed (unpaired) axes. As homologs synapse, ATR disappeared from these locations. However, at regions delayed in synapsis, often seen near the proximal ends of autosomal bivalents, there was an accumulation of ATR foci along the unsynapsed axes. ATR was detected at similar locations on the two axial elements. In nuclei where an entire autosome fails to find its homologous pairing partner, ATR foci were detected along the entire lengths of these asynapsed axis. In males, where the X chromosome has no homolog, ATR foci were localized along the unpaired axis.

ATM was also visualized as foci and was first detected during zygonema as homologs synapse, but ATM localization was different than ATR. ATM was first observed along synapsed axes when homologous autosomal axial elements come into contact. However, during mid-pachynema, after autosomal synapsis has been completed, ATM foci appeared on the X chromosome axis. ATM localization persisted on fully synapsed bivalents into pachynema, a substage that lasts 3 days in mouse oocytes and 6 days in mouse spermatocytes. During pachynema, the number of foci drops gradually, stabilizing briefly in mid-pachynema before eventually disappearing mid-to late pachynema. Thus, ATR and ATM protein kinases play important and complementary roles at distinct stages in meiosis I.

The involvement of ATR appears to be transient during early meiotic prophase while the role of ATM appears to be more prolonged. However, both ATR and ATM coordinate the various events of meiotic prophase by performing similar checkpoint functions.

The foregoing illustrative examples relate to presently preferred embodiments of the invention and numerous modifications and variations thereof are expected to occur to those skilled in the art. Thus only such limitations as appear in the appended claims should be placed upon the scope of the present invention.



-51-

## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

- (i) APPLICANT: ICOS Corporation
- (ii) TITLE OF INVENTION: Cell Cycle Checkpoint PIK-Related Kinase  
Materials and Methods
- (iii) NUMBER OF SEQUENCES: 42
- (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
  - (B) STREET: 6300 Sears Tower, 233 S. Wacker Dr.
  - (C) CITY: Chicago
  - (D) STATE: Illinois
  - (E) COUNTRY: USA
  - (F) ZIP: 60606
- (v) COMPUTER READABLE FORM:
  - (A) MEDIUM TYPE: Floppy disk
  - (B) COMPUTER: IBM PC compatible
  - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
  - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
- (vi) CURRENT APPLICATION DATA:
  - (A) APPLICATION NUMBER:
  - (B) FILING DATE:
  - (C) CLASSIFICATION:
- (viii) ATTORNEY/AGENT INFORMATION:
  - (A) NAME: Noland, Greta E.
  - (B) REGISTRATION NUMBER: 35,302
  - (C) REFERENCE/DOCKET NUMBER: 27866/33607
- (ix) TELECOMMUNICATION INFORMATION:
  - (A) TELEPHONE: (312) 474-6300
  - (B) TELEFAX: (312) 474-0448
  - (C) TELEX: 25-3856

## (2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 7621 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: pBSHFB2HT2-27
- (ix) FEATURE:
  - (A) NAME/KEY: CDS
  - (B) LOCATION: 333..7559
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

-52-

CTTGTGAAGA	GAATGTTTTA	CACTCTTGTT	AGTGAAGTTT	ATTCTTTAAA	AGTCAATCGT	60										
CAAGGATTTA	GCAAATGAAT	TAGCACTTCG	GATATACTTG	TTTATTTAAT	ATCTTTTTTTG	120										
TTTATTTCAA	AGAATTCAGT	AATTGGATCA	TAACGAGACT	TCTGCGGATT	GCAGCAACTC	180										
CCTCCTGTCA	TTTGTTACAC	AAGAAAATCT	GTGAAGTCAT	CTGTTTCATTA	TTATTTCTTT	240										
TTAAAAGCAA	GAGTCCTGCT	ATTTTTGGGG	TACTCACAAA	AGAATTATTA	CAACTTTTTTG	300										
AAGACTTGGT	TTACCTCCAT	AGAAGAAATG	TG	ATG	GGT	CAT	GCT	GTG	GAA	TGG	353					
				Met	Gly	His	Ala	Val	Glu	Trp						
				1				5								
CCA	GTG	GTC	ATG	AGC	CGA	TTT	TTA	AGT	CAA	TTA	GAT	GAA	CAC	ATG	GGA	401
Pro	Val	Val	Met	Ser	Arg	Phe	Leu	Ser	Gln	Leu	Asp	Glu	His	Met	Gly	
	10						15					20				
TAT	TTA	CAA	TCA	GCT	CCT	TTG	CAG	TTG	ATG	AGT	ATG	CAA	AAA	TTA	GAA	449
Tyr	Leu	Gln	Ser	Ala	Pro	Leu	Gln	Leu	Met	Ser	Met	Gln	Lys	Leu	Glu	
	25					30					35					
TTT	ATT	GAA	GTC	ACT	TTA	TTA	ACG	GTT	CTT	ACT	CGT	ATT	ATT	GCA	ATT	497
Phe	Ile	Glu	Val	Thr	Leu	Leu	Thr	Val	Leu	Thr	Arg	Ile	Ile	Ala	Ile	
	40				45					50					55	
GTG	TTT	TTT	AGA	AGG	CAA	GAA	CTC	TTA	CTT	TGG	CAG	ATA	GGT	TGT	GTT	545
Val	Phe	Phe	Arg	Arg	Gln	Glu	Leu	Leu	Leu	Trp	Gln	Ile	Gly	Cys	Val	
				60					65					70		
CTG	CTA	GAG	TAT	GGT	AGT	CCA	AAA	ATT	AAA	TCC	CTA	GCA	ATT	AGC	TTT	593
Leu	Leu	Glu	Tyr	Gly	Ser	Pro	Lys	Ile	Lys	Ser	Leu	Ala	Ile	Ser	Phe	
			75					80						85		
TTA	ACA	GAA	CTT	TTT	CAG	CTT	GGA	GGA	CTA	CCA	GCA	CAA	CCA	GCT	AGC	641
Leu	Thr	Glu	Leu	Phe	Gln	Leu	Gly	Gly	Leu	Pro	Ala	Gln	Pro	Ala	Ser	
	90						95					100				
ACT	TTT	TTC	AGC	TCA	TTT	TTG	GAA	TTA	TTA	AAA	CAC	CTT	GTA	GAA	ATG	689
Thr	Phe	Phe	Ser	Ser	Phe	Leu	Glu	Leu	Leu	Lys	His	Leu	Val	Glu	Met	
	105					110					115					
GAT	ACT	GAC	CAA	TTG	AAA	CTC	TAT	GAA	GAG	CCA	TTA	TCA	AAG	CTG	ATA	737
Asp	Thr	Asp	Gln	Leu	Lys	Leu	Tyr	Glu	Glu	Pro	Leu	Ser	Lys	Leu	Ile	
	120				125					130					135	
AAG	ACA	CTA	TTT	CCC	TTT	GAA	GCA	GAA	GCT	TAT	AGA	AAT	ATT	GAA	CCT	785
Lys	Thr	Leu	Phe	Pro	Phe	Glu	Ala	Glu	Ala	Tyr	Arg	Asn	Ile	Glu	Pro	
				140					145					150		
GTC	TAT	TTA	AAT	ATG	CTG	CTG	GAA	AAA	CTC	TGT	GTC	ATG	TTT	GAA	GAC	833
Val	Tyr	Leu	Asn	Met	Leu	Leu	Glu	Lys	Leu	Cys	Val	Met	Phe	Glu	Asp	
			155					160						165		
GGT	GTG	CTC	ATG	CGG	CTT	AAG	TCT	GAT	TTG	CTA	AAA	GCA	GCT	TTG	TGC	881
Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	Leu	Leu	Lys	Ala	Ala	Leu	Cys	
		170					175					180				
CAT	TTA	CTG	CAG	TAT	TTC	CTT	AAA	TTT	GTG	CCA	GCT	GGG	TAT	GAA	TCT	929
His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	Val	Pro	Ala	Gly	Tyr	Glu	Ser	
	185					190					195					
GCT	TTA	CAA	GTC	AGG	AAG	GTC	TAT	GTG	AGA	AAT	ATT	TGT	AAA	GCT	CTT	977
Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	Arg	Asn	Ile	Cys	Lys	Ala	Leu	
	200				205					210					215	

-53-

TTG	GAT	GTG	CTT	GGA	ATT	GAG	GTA	GAT	GCA	GAG	TAC	TTG	TTG	GGC	CCA	1025
Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	Ala	Glu	Tyr	Leu	Leu	Gly	Pro	
				220					225					230		
CTT	TAT	GCA	GCT	TTG	AAA	ATG	GAA	AGT	ATG	GAA	ATC	ATT	GAG	GAG	ATT	1073
Leu	Tyr	Ala	Ala	Leu	Lys	Met	Glu	Ser	Met	Glu	Ile	Ile	Glu	Glu	Ile	
			235					240					245			
CAA	TGC	CAA	ACT	CAA	CAG	GAA	AAC	CTC	AGC	AGT	AAT	AGT	GAT	GGA	ATA	1121
Gln	Cys	Gln	Thr	Gln	Gln	Glu	Asn	Leu	Ser	Ser	Asn	Ser	Asp	Gly	Ile	
		250					255					260				
TCA	CCC	AAA	AGG	CGT	CGT	CTC	AGC	TCG	TCT	CTA	AAC	CCT	TCT	AAA	AGA	1169
Ser	Pro	Lys	Arg	Arg	Arg	Leu	Ser	Ser	Ser	Leu	Asn	Pro	Ser	Lys	Arg	
	265					270					275					
GCA	CCA	AAA	CAG	ACT	GAG	GAA	ATT	AAA	CAT	GTG	GAC	ATG	AAC	CAA	AAG	1217
Ala	Pro	Lys	Gln	Thr	Glu	Glu	Ile	Lys	His	Val	Asp	Met	Asn	Gln	Lys	
	280				285					290					295	
AGC	ATA	TTA	TGG	AGT	GCA	CTG	AAA	CAG	AAA	GCT	GAA	TCC	CTT	CAG	ATT	1265
Ser	Ile	Leu	Trp	Ser	Ala	Leu	Lys	Gln	Lys	Ala	Glu	Ser	Leu	Gln	Ile	
				300					305					310		
TCC	CTT	GAA	TAC	AGT	GGC	CTA	AAG	AAT	CCT	GTT	ATT	GAG	ATG	TTA	GAA	1313
Ser	Leu	Glu	Tyr	Ser	Gly	Leu	Lys	Asn	Pro	Val	Ile	Glu	Met	Leu	Glu	
			315					320					325			
GGA	ATT	GCT	GTT	GTC	TTA	CAA	CTG	ACT	GCT	CTG	TGT	ACT	GTT	CAT	TGT	1361
Gly	Ile	Ala	Val	Val	Leu	Gln	Leu	Thr	Ala	Leu	Cys	Thr	Val	His	Cys	
		330					335					340				
TCT	CAT	CAA	AAC	ATG	AAC	TGC	CGT	ACT	TTC	AAG	GAC	TGT	CAA	CAT	AAA	1409
Ser	His	Gln	Asn	Met	Asn	Cys	Arg	Thr	Phe	Lys	Asp	Cys	Gln	His	Lys	
	345					350					355					
TCC	AAG	AAG	AAA	CCT	TCT	GTA	GTG	ATA	ACT	TGG	ATG	TCA	TTG	GAT	TTT	1457
Ser	Lys	Lys	Lys	Pro	Ser	Val	Val	Ile	Thr	Trp	Met	Ser	Leu	Asp	Phe	
	360					365				370					375	
TAC	ACA	ACA	GTG	CTT	AAG	AGC	TGT	AGA	AGG	TTG	TTA	GAA	TCT	GTT	CAG	1505
Tyr	Thr	Thr	Val	Leu	Lys	Ser	Cys	Arg	Arg	Leu	Leu	Glu	Ser	Val	Gln	
				380					385					390		
AAA	CGG	ACT	GGA	GGC	AAC	ATT	GAT	AAG	GTG	GTG	AAA	ATT	TAT	GAT	GCT	1553
Lys	Arg	Thr	Gly	Gly	Asn	Ile	Asp	Lys	Val	Val	Lys	Ile	Tyr	Asp	Ala	
			395					400					405			
TTG	ATT	TAT	ATG	CAA	GTA	AAC	AGT	TCA	TTT	GAA	GAT	CAT	ATC	CTG	GAA	1601
Leu	Ile	Tyr	Met	Gln	Val	Asn	Ser	Ser	Phe	Glu	Asp	His	Ile	Leu	Glu	
		410					415					420				
GAT	TTA	TGT	GGA	ATG	CTC	TCA	CTT	CCA	TGG	ATT	TAT	TCC	CAT	TCT	GAT	1649
Asp	Leu	Cys	Gly	Met	Leu	Ser	Leu	Pro	Trp	Ile	Tyr	Ser	His	Ser	Asp	
	425					430					435					
GAT	GGC	TGT	TTA	AAG	TTG	ACC	ACA	TTT	GCC	GCT	AAT	CTT	CTA	ACA	TTA	1697
Asp	Gly	Cys	Leu	Lys	Leu	Thr	Thr	Phe	Ala	Ala	Asn	Leu	Leu	Thr	Leu	
	440				445				450						455	
AGC	TGT	AGG	ATT	TCA	GAT	AGC	TAT	TCA	CCA	CAG	GCA	CAA	TCA	CGA	TGT	1745
Ser	Cys	Arg	Ile	Ser	Asp	Ser	Tyr	Ser	Pro	Gln	Ala	Gln	Ser	Arg	Cys	
				460					465					470		
GTG	TTT	CTT	CTG	ACT	CTG	TTT	CCA	AGA	AGA	ATA	TTC	CTT	GAG	TGG	AGA	1793
Val	Phe	Leu	Leu	Thr	Leu	Phe	Pro	Arg	Arg	Ile	Phe	Leu	Glu	Trp	Arg	
			475					480					485			

-54-

ACA	GCA	GTT	TAC	AAC	TGG	GCC	CTG	CAG	AGC	TCC	CAT	GAA	GTA	ATC	CGG	1841
Thr	Ala	Val	Tyr	Asn	Trp	Ala	Leu	Gln	Ser	Ser	His	Glu	Val	Ile	Arg	
		490					495					500				
GCT	AGT	TGT	GTT	AGT	GGA	TTT	TTT	ATC	TTA	TTG	CAG	CAG	CAG	AAT	TCT	1889
Ala	Ser	Cys	Val	Ser	Gly	Phe	Phe	Ile	Leu	Leu	Gln	Gln	Gln	Asn	Ser	
	505					510					515					
TGT	AAC	AGA	GTT	CCC	AAG	ATT	CTT	ATA	GAT	AAA	GTC	AAA	GAT	GAT	TCT	1937
Cys	Asn	Arg	Val	Pro	Lys	Ile	Leu	Ile	Asp	Lys	Val	Lys	Asp	Asp	Ser	
520					525					530					535	
GAC	ATT	GTC	AAG	AAA	GAA	TTT	GCT	TCT	ATA	CTT	GGT	CAA	CTT	GTC	TGT	1985
Asp	Ile	Val	Lys	Lys	Glu	Phe	Ala	Ser	Ile	Leu	Gly	Gln	Leu	Val	Cys	
				540				545						550		
ACT	CTT	CAC	GGC	ATG	TTT	TAT	CTG	ACA	AGT	TCT	TTA	ACA	GAA	CCT	TTC	2033
Thr	Leu	His	Gly	Met	Phe	Tyr	Leu	Thr	Ser	Ser	Leu	Thr	Glu	Pro	Phe	
			555					560					565			
TCT	GAA	CAC	GGA	CAT	GTG	GAC	CTC	TTC	TGT	AGG	AAC	TTG	AAA	GCC	ACT	2081
Ser	Glu	His	Gly	His	Val	Asp	Leu	Phe	Cys	Arg	Asn	Leu	Lys	Ala	Thr	
		570					575					580				
TCT	CAA	CAT	GAA	TGT	TCA	TCT	TCT	CAA	CTA	AAA	GCT	TCT	GTC	TGC	AAG	2129
Ser	Gln	His	Glu	Cys	Ser	Ser	Ser	Gln	Leu	Lys	Ala	Ser	Val	Cys	Lys	
	585					590					595					
CCA	TTC	CTT	TTC	CTA	CTG	AAA	AAA	AAA	ATA	CCT	AGT	CCA	GTA	AAA	CTT	2177
Pro	Phe	Leu	Phe	Leu	Leu	Lys	Lys	Lys	Ile	Pro	Ser	Pro	Val	Lys	Leu	
600					605					610					615	
GCT	TTC	ATA	GAT	AAT	CTA	CAT	CAT	CTT	TGT	AAG	CAT	CTT	GAT	TTT	AGA	2225
Ala	Phe	Ile	Asp	Asn	Leu	His	His	Leu	Cys	Lys	His	Leu	Asp	Phe	Arg	
				620					625					630		
GAA	GAT	GAA	ACA	GAT	GTA	AAA	GCA	GTT	CTT	GGA	ACT	TTA	TTA	AAT	TTA	2273
Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala	Val	Leu	Gly	Thr	Leu	Leu	Asn	Leu	
			635					640					645			
ATG	GAA	GAT	CCA	GAC	AAA	GAT	GTT	AGA	GTG	GCT	TTT	AGT	GGA	AAT	ATC	2321
Met	Glu	Asp	Pro	Asp	Lys	Asp	Val	Arg	Val	Ala	Phe	Ser	Gly	Asn	Ile	
		650				655						660				
AAG	CAC	ATA	TTG	GAA	TCC	TTG	GAC	TCT	GAA	GAT	GGA	TTT	ATA	AAG	GAG	2369
Lys	His	Ile	Leu	Glu	Ser	Leu	Asp	Ser	Glu	Asp	Gly	Phe	Ile	Lys	Glu	
	665					670					675					
CTT	TTT	GTC	TTA	AGA	ATG	AAG	GAA	GCA	TAT	ACA	CAT	GCC	CAA	ATA	TCA	2417
Leu	Phe	Val	Leu	Arg	Met	Lys	Glu	Ala	Tyr	Thr	His	Ala	Gln	Ile	Ser	
680					685					690					695	
AGA	AAT	AAT	GAG	CTG	AAG	GAT	ACC	TTG	ATT	CTT	ACA	ACA	GGG	GAT	ATT	2465
Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr	Leu	Ile	Leu	Thr	Thr	Gly	Asp	Ile	
				700				705						710		
GGA	AGG	GCC	GCA	AAA	GGA	GAT	TTG	GTA	CCA	TTT	GCA	CTC	TTA	CAC	TTA	2513
Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu	Val	Pro	Phe	Ala	Leu	Leu	His	Leu	
			715					720					725			
TTG	CAT	TGT	TTG	TTA	TCC	AAG	TCA	GCA	TCT	GTC	TCT	GGA	GCA	GCA	TAC	2561
Leu	His	Cys	Leu	Leu	Ser	Lys	Ser	Ala	Ser	Val	Ser	Gly	Ala	Ala	Tyr	
		730				735						740				
ACA	GAA	ATT	AGA	GCT	CTG	GTT	GCA	GCT	AAA	AGT	GTT	AAA	CTG	CAA	AGT	2609
Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala	Ala	Lys	Ser	Val	Lys	Leu	Gln	Ser	
	745					750					755					

-55-

TTT	TTC	AGC	CAG	TAT	AAG	AAA	CCC	ATC	TGT	CAG	TTT	TTG	GTA	GAA	TCC	2657
Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro	Ile	Cys	Gln	Phe	Leu	Val	Glu	Ser	
760					765				770						775	
CTT	CAC	TCT	AGT	CAG	ATG	ACA	GCA	CTT	CCG	AAT	ACT	CCA	TGC	CAG	AAT	2705
Leu	His	Ser	Ser	Gln	Met	Thr	Ala	Leu	Pro	Asn	Thr	Pro	Cys	Gln	Asn	
				780					785						790	
GCT	GAC	GTG	CGA	AAA	CAA	GAT	GTG	GCT	CAC	CAG	AGA	GAA	ATG	GCT	TTA	2753
Ala	Asp	Val	Arg	Lys	Gln	Asp	Val	Ala	His	Gln	Arg	Glu	Met	Ala	Leu	
			795					800					805			
AAT	ACG	TTG	TCT	GAA	ATT	GCC	AAC	GTT	TTC	GAC	TTT	CCT	GAT	CTT	AAT	2801
Asn	Thr	Leu	Ser	Glu	Ile	Ala	Asn	Val	Phe	Asp	Phe	Pro	Asp	Leu	Asn	
		810					815					820				
CGT	TTT	CTT	ACT	AGG	ACA	TTA	CAA	GTT	CTA	CTA	CCT	GAT	CTT	GCT	GCC	2849
Arg	Phe	Leu	Thr	Arg	Thr	Leu	Gln	Val	Leu	Leu	Pro	Asp	Leu	Ala	Ala	
	825					830					835					
AAA	GCA	AGC	CCT	GCA	GCT	TCT	GCT	CTC	ATT	CGA	ACT	TTA	GGA	AAA	CAA	2897
Lys	Ala	Ser	Pro	Ala	Ala	Ser	Ala	Leu	Ile	Arg	Thr	Leu	Gly	Lys	Gln	
840					845					850					855	
TTA	AAT	GTC	AAT	CGT	AGA	GAG	ATT	TTA	ATA	AAC	AAC	TTC	AAA	TAT	ATT	2945
Leu	Asn	Val	Asn	Arg	Arg	Glu	Ile	Leu	Ile	Asn	Asn	Phe	Lys	Tyr	Ile	
				860					865					870		
TTT	TCT	CAT	TTG	GTC	TGT	TCT	TGT	TCC	AAA	GAT	GAA	TTA	GAA	CGT	GCC	2993
Phe	Ser	His	Leu	Val	Cys	Ser	Cys	Ser	Lys	Asp	Glu	Leu	Glu	Arg	Ala	
			875					880					885			
CTT	CAT	TAT	CTG	AAG	AAT	GAA	ACA	GAA	ATT	GAA	CTG	GGG	AGC	CTG	TTG	3041
Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr	Glu	Ile	Glu	Leu	Gly	Ser	Leu	Leu	
		890					895					900				
AGA	CAA	GAT	TTC	CAA	GGA	TTG	CAT	AAT	GAA	TTA	TTG	CTG	CGT	ATT	GGA	3089
Arg	Gln	Asp	Phe	Gln	Gly	Leu	His	Asn	Glu	Leu	Leu	Leu	Arg	Ile	Gly	
	905					910					915					
GAA	CAC	TAT	CAA	CAG	GTT	TTT	AAT	GGT	TTG	TCA	ATA	CTT	GCC	TCA	TTT	3137
Glu	His	Tyr	Gln	Gln	Val	Phe	Asn	Gly	Leu	Ser	Ile	Leu	Ala	Ser	Phe	
920					925				930						935	
GCA	TCC	AGT	GAT	GAT	CCA	TAT	CAG	GGC	CCG	AGA	GAT	ATC	ATA	TCA	CCT	3185
Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln	Gly	Pro	Arg	Asp	Ile	Ile	Ser	Pro	
				940					945					950		
GAA	CTG	ATG	GCT	GAT	TAT	TTA	CAA	CCC	AAA	TTG	TTG	GGC	ATT	TTG	GCT	3233
Glu	Leu	Met	Ala	Asp	Tyr	Leu	Gln	Pro	Lys	Leu	Leu	Gly	Ile	Leu	Ala	
			955					960					965			
TTT	TTT	AAC	ATG	CAG	TTA	CTG	AGC	TCT	AGT	GTT	GGC	ATT	GAA	GAT	AAG	3281
Phe	Phe	Asn	Met	Gln	Leu	Leu	Ser	Ser	Ser	Val	Gly	Ile	Glu	Asp	Lys	
		970					975					980				
AAA	ATG	GCC	TTG	AAC	AGT	TTG	ATG	TCT	TTG	ATG	AAG	TTA	ATG	GGA	CCC	3329
Lys	Met	Ala	Leu	Asn	Ser	Leu	Met	Ser	Leu	Met	Lys	Leu	Met	Gly	Pro	
	985					990					995					
AAA	CAT	GTC	AGT	TCT	GTG	AGG	GTG	AAG	ATG	ATG	ACC	ACA	CTG	AGA	ACT	3377
Lys	His	Val	Ser	Ser	Val	Arg	Val	Lys	Met	Met	Thr	Thr	Leu	Arg	Thr	
1000					1005				1010						1015	
GGC	CTT	CGA	TTC	AAG	GAT	GAT	TTT	CCT	GAA	TTG	TGT	TGC	AGA	GCT	TGG	3425
Gly	Leu	Arg	Phe	Lys	Asp	Asp	Phe	Pro	Glu	Leu	Cys	Cys	Arg	Ala	Trp	
				1020					1025					1030		

-56-

GAC TGC TTT GTT CGC TGC CTG GAT CAT GCT TGT CTG GGC TCC CTT CTC	3473
Asp Cys Phe Val Arg Cys Leu Asp His Ala Cys Leu Gly Ser Leu Leu	
1035 1040 1045	
AGT CAT GTA ATA GTA GCT TTG TTA CCT CTT ATA CAC ATC CAG CCT AAA	3521
Ser His Val Ile Val Ala Leu Leu Pro Leu Ile His Ile Gln Pro Lys	
1050 1055 1060	
GAA ACT GCA GCT ATC TTC CAC TAC CTC ATA ATT GAA AAC AGG GAT GCT	3569
Glu Thr Ala Ala Ile Phe His Tyr Leu Ile Ile Glu Asn Arg Asp Ala	
1065 1070 1075	
GTG CAA GAT TTT CTT CAT GAA ATA TAT TTT TTA CCT GAT CAT CCA GAA	3617
Val Gln Asp Phe Leu His Glu Ile Tyr Phe Leu Pro Asp His Pro Glu	
1080 1085 1090 1095	
TTA AAA AAG ATA AAA GCC GTT CTC CAG GAA TAC AGA AAG GAG ACC TCT	3665
Leu Lys Lys Ile Lys Ala Val Leu Gln Glu Tyr Arg Lys Glu Thr Ser	
1100 1105 1110	
GAG AGC ACT GAT CTT CAG ACA ACT CTT CAG CTC TCT ATG AAG GCC ATT	3713
Glu Ser Thr Asp Leu Gln Thr Thr Leu Gln Leu Ser Met Lys Ala Ile	
1115 1120 1125	
CAA CAT GAA AAT GTC GAT GTT CGT ATT CAT GCT CTT ACA AGC TTG AAG	3761
Gln His Glu Asn Val Asp Val Arg Ile His Ala Leu Thr Ser Leu Lys	
1130 1135 1140	
GAA ACC TTG TAT AAA AAT CAG GAA AAA CTG ATA AAG TAT GCA ACA GAC	3809
Glu Thr Leu Tyr Lys Asn Gln Glu Lys Leu Ile Lys Tyr Ala Thr Asp	
1145 1150 1155	
AGT GAA ACA GTA GAA CCT ATT ATC TCA CAG TTG GTG ACA GTG CTT TTG	3857
Ser Glu Thr Val Glu Pro Ile Ile Ser Gln Leu Val Thr Val Leu Leu	
1160 1165 1170 1175	
AAA GGT TGC CAA GAT GCA AAC TCT CAA GCT CGG TTG CTC TGT GGG GAA	3905
Lys Gly Cys Gln Asp Ala Asn Ser Gln Ala Arg Leu Leu Cys Gly Glu	
1180 1185 1190	
TGT TTA GGG GAA TTG GGG GCG ATA GAT CCA GGT CGA TTA GAT TTC TCA	3953
Cys Leu Gly Glu Leu Gly Ala Ile Asp Pro Gly Arg Leu Asp Phe Ser	
1195 1200 1205	
ACA ACT GAA ACT CAA GGA AAA GAT TTT ACA TTT GTG ACT GGA GTA GAA	4001
Thr Thr Glu Thr Gln Gly Lys Asp Phe Thr Phe Val Thr Gly Val Glu	
1210 1215 1220	
GAT TCA AGC TTT GCC TAT GGA TTA TTG ATG GAG CTA ACA AGA GCT TAC	4049
Asp Ser Ser Phe Ala Tyr Gly Leu Leu Met Glu Leu Thr Arg Ala Tyr	
1225 1230 1235	
CTT GCG TAT GCT GAT AAT AGC CGA GCT CCA GAT TCA GCT GCC TAT GCC	4097
Leu Ala Tyr Ala Asp Asn Ser Arg Ala Pro Asp Ser Ala Ala Tyr Ala	
1240 1245 1250 1255	
ATT CAG GAG TTG CTT TCT ATT TAT GAC TGT AGA GAG ATG GAG ACC AAC	4145
Ile Gln Glu Leu Leu Ser Ile Tyr Asp Cys Arg Glu Met Glu Thr Asn	
1260 1265 1270	
GGC CCA GGT CAC CAA TTG TGG AGG AGA TTT CCT GAG CAT GTT CGG GAA	4193
Gly Pro Gly His Gln Leu Trp Arg Arg Phe Pro Glu His Val Arg Glu	
1275 1280 1285	
ATA CTA GAA CCT CAT CTA AAT ACC AGA TAC AAG AGT TCT CAG AAG TCA	4241
Ile Leu Glu Pro His Leu Asn Thr Arg Tyr Lys Ser S r Gln Lys Ser	
1290 1295 1300	

-57-

ACC GAT TGG TCT GGA GTA AAG AAG CCA ATT TAC TTA AGT AAA TTG GGT Thr Asp Trp Ser Gly Val Lys Lys Pro Ile Tyr Leu Ser Lys Leu Gly 1305 1310 1315	4289
AGT AAC TTT GCA GAA TGG TCA GCA TCT TGG GCA GGT TAT CTT ATT ACA Ser Asn Phe Ala Glu Trp Ser Ala Ser Trp Ala Gly Tyr Leu Ile Thr 1320 1325 1330 1335	4337
AAG GTT CGA CAT GAT CTT GCC AGT AAA ATT TTC ACC TGC TGT AGC ATT Lys Val Arg His Asp Leu Ala Ser Lys Ile Phe Thr Cys Cys Ser Ile 1340 1345 1350	4385
ATG ATG AAG CAT GAT TTC AAA GTG ACC ATC TAT CTT CTT CCA CAT ATT Met Met Lys His Asp Phe Lys Val Thr Ile Tyr Leu Leu Pro His Ile 1355 1360 1365	4433
CTG GTG TAT GTC TTA CTG GGT TGT AAT CAA GAA GAT CAG CAG GAG GTT Leu Val Tyr Val Leu Leu Gly Cys Asn Gln Glu Asp Gln Gln Glu Val 1370 1375 1380	4481
TAT GCA GAA ATT ATG GCA GTT CTA AAG CAT GAC GAT CAG CAT ACC ATA Tyr Ala Glu Ile Met Ala Val Leu Lys His Asp Asp Gln His Thr Ile 1385 1390 1395	4529
AAT ACC CAA GAC ATT GCA TCT GAT CTG TGT CAA CTC AGT ACA CAG ACT Asn Thr Gln Asp Ile Ala Ser Asp Leu Cys Gln Leu Ser Thr Gln Thr 1400 1405 1410 1415	4577
GTG TTC TCC ATG CTT GAC CAT CTC ACA CAG TGG GCA AGG CAC AAA TTT Val Phe Ser Met Leu Asp His Leu Thr Gln Trp Ala Arg His Lys Phe 1420 1425 1430	4625
CAG GCA CTG AAA GCT GAG AAA TGT CCA CAC AGC AAA TCA AAC AGA AAT Gln Ala Leu Lys Ala Glu Lys Cys Pro His Ser Lys Ser Asn Arg Asn 1435 1440 1445	4673
AAG GTA GAC TCA ATG GTA TCT ACT GTG GAT TAT GAA GAC TAT CAG AGT Lys Val Asp Ser Met Val Ser Thr Val Asp Tyr Glu Asp Tyr Gln Ser 1450 1455 1460	4721
GTA ACC CGT TTT CTA GAC CTC ATA CCC CAG GAT ACT CTG GCA GTA GCT Val Thr Arg Phe Leu Asp Leu Ile Pro Gln Asp Thr Leu Ala Val Ala 1465 1470 1475	4769
TCC TTT CGC TCC AAA GCA TAC ACA CGA GCT GTA ATG CAC TTT GAA TCA Ser Phe Arg Ser Lys Ala Tyr Thr Arg Ala Val Met His Phe Glu Ser 1480 1485 1490 1495	4817
TTT ATT ACA GAA AAG AAG CAA AAT ATT CAG GAA CAT CTT GGA TTT TTA Phe Ile Thr Glu Lys Lys Gln Asn Ile Gln Glu His Leu Gly Phe Leu 1500 1505 1510	4865
CAG AAA TTG TAT GCT GCT ATG CAT GAA CCT GAT GGA GTG TCC GGA GTC Gln Lys Leu Tyr Ala Ala Met His Glu Pro Asp Gly Val Ser Gly Val 1515 1520 1525	4913
AGT GCA ATT AGA AAG GCA GAA CCA TCT CTA AAA GAA CAG ATC CTT GAA Ser Ala Ile Arg Lys Ala Glu Pro Ser Leu Lys Glu Gln Ile Leu Glu 1530 1535 1540	4961
CAT GAA AGC CTT GGC TTG CTG AGG GAT GCC ACT GCT TGT TAT GAC AGG His Glu Ser Leu Gly Leu Leu Arg Asp Ala Thr Ala Cys Tyr Asp Arg 1545 1550 1555	5009
GCT ATT CAG CTA GAA CCA GAC CAG ATC ATT CAT TAC CAT GGT GTA GTA Ala Ile Gln Leu Glu Pro Asp Gln Ile Ile His Tyr His Gly Val Val 1560 1565 1570 1575	5057

-58-

AAG TCC ATG TTA GGT CTT GGT CAG CTG TCT ACT GTT ATC ACT CAG GTG Lys Ser Met Leu Gly Leu Gly Gln Leu Ser Thr Val Ile Thr Gln Val 1580 1585 1590	5105
AAT GGA GTG CAT GCT AAC AGG TCC GAG TGG ACA GAT GAA TTA AAC ACG Asn Gly Val His Ala Asn Arg Ser Glu Trp Thr Asp Glu Leu Asn Thr 1595 1600 1605	5153
TAC AGA GTG GAA GCA GCT TGG AAA TTG TCA CAG TGG GAT TTG GTG GAA Tyr Arg Val Glu Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val Glu 1610 1615 1620	5201
AAC TAT TTG GCA GCA GAT GGA AAA TCT ACA ACA TGG AGT GTC AGA CTG Asn Tyr Leu Ala Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg Leu 1625 1630 1635	5249
GGA CAG CTA TTA TTA TCA GCC AAA AAA AGA GAT ATC ACA GCT TTT TAT Gly Gln Leu Leu Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe Tyr 1640 1645 1650 1655	5297
GAC TCA CTG AAA CTA GTG AGA GCA GAA CAA ATT GTA CCT CTT TCA GCT Asp Ser Leu Lys Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser Ala 1660 1665 1670	5345
GCA AGC TTT GAA AGA GGC TCC TAC CAA CGA GGA TAT GAA TAT ATT GTG Ala Ser Phe Glu Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile Val 1675 1680 1685	5393
AGA TTG CAC ATG TTA TGT GAG TTG GAG CAT AGC ATC AAA CCA CTT TTC Arg Leu His Met Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu Phe 1690 1695 1700	5441
CAG CAT TCT CCA GGT GAC AGT TCT CAA GAA GAT TCT CTA AAC TGG GTA Gln His Ser Pro Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp Val 1705 1710 1715	5489
GCT CGA CTA GAA ATG ACC CAG AAT TCC TAC AGA GCC AAG GAG CCT ATC Ala Arg Leu Glu Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro Ile 1720 1725 1730 1735	5537
CTG GCT CTC CGG AGG GCT TTA CTA AGC CTC AAC AAA AGA CCA GAT TAC Leu Ala Leu Arg Arg Ala Leu Leu Ser Leu Asn Lys Arg Pro Asp Tyr 1740 1745 1750	5585
AAT GAA ATG GTT GGA GAA TGC TGG CTG CAG AGT GCC AGG GTA GCT AGA Asn Glu Met Val Gly Glu Cys Trp Leu Gln Ser Ala Arg Val Ala Arg 1755 1760 1765	5633
AAG GCT GGT CAC CAC CAG ACA GCC TAC AAT GCT CTC CTT AAT GCA GGG Lys Ala Gly His His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala Gly 1770 1775 1780	5681
GAA TCA CGA CTC GCT GAA CTG TAC GTG GAA AGG GCA AAG TGG CTC TGG Glu Ser Arg Leu Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu Trp 1785 1790 1795	5729
TCC AAG GGT GAT GTT CAC CAG GCA CTA ATT GTT CTT CAA AAA GGT GTT Ser Lys Gly Asp Val His Gln Ala Leu Ile Val Leu Gln Lys Gly Val 1800 1805 1810 1815	5777
GAA TTA TGT TTT CCT GAA AAT GAA ACC CCA CCT GAG GGT AAG AAC ATG Glu Leu Cys Phe Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn Met 1820 1825 1830	5825
TTA ATC CAT GGT CGA GCT ATG CTA CTA GTG GGC CGA TTT ATG GAA GAA Leu Ile His Gly Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu Glu 1835 1840 1845	5873



-59-

ACA GCT AAC TTT GAA AGC AAT GCA ATT ATG AAA AAA TAT AAG GAT GTG	5921
Thr Ala Asn Phe Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp Val	
1850 1855 1860	
ACC GCG TGC CTG CCA GAA TGG GAG GAT GGG CAT TTT TAC CTT GCC AAG	5969
Thr Ala Cys Leu Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala Lys	
1865 1870 1875	
TAC TAT GAC AAA TTG ATG CCC ATG GTC ACA GAC AAC AAA ATG GAA AAG	6017
Tyr Tyr Asp Lys Leu Met Pro Met Val Thr Asp Asn Lys Met Glu Lys	
1880 1885 1890 1895	
CAA GGT GAT CTC ATC CGG TAT ATA GTT CTT CAT TTT GGC AGA TCT CTA	6065
Gln Gly Asp Leu Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser Leu	
1900 1905 1910	
CAA TAT GGA AAT CAG TTC ATA TAT CAG TCA ATG CCA CGA ATG TTA ACT	6113
Gln Tyr Gly Asn Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu Thr	
1915 1920 1925	
CTA TGG CTT GAT TAT GGT ACA AAG GCA TAT GAA TGG GAA AAA GCT GGC	6161
Leu Trp Leu Asp Tyr Gly Thr Lys Ala Tyr Glu Trp Glu Lys Ala Gly	
1930 1935 1940	
CGC TCC GAT CGT GTA CAA ATG AGG AAT GAT TTG GGT AAA ATA AAC AAG	6209
Arg Ser Asp Arg Val Gln Met Arg Asn Asp Leu Gly Lys Ile Asn Lys	
1945 1950 1955	
GTT ATC ACA GAG CAT ACA AAC TAT TTA GCT CCA TAT CAA TTT TTG ACT	6257
Val Ile Thr Glu His Thr Asn Tyr Leu Ala Pro Tyr Gln Phe Leu Thr	
1960 1965 1970 1975	
GCT TTT TCA CAA TTG ATC TCT CGA ATT TGT CAT TCT CAC GAT GAA GTT	6305
Ala Phe Ser Gln Leu Ile Ser Arg Ile Cys His Ser His Asp Glu Val	
1980 1985 1990	
TTT GTT GTG CTT GAT GGA AAT AAT AGC CAA GTA TTT CTA GCC TAT CCT	6353
Phe Val Val Leu Asp Gly Asn Asn Ser Gln Val Phe Leu Ala Tyr Pro	
1995 2000 2005	
CAA CAA GCA ATG TGG ATG ATG ACA GCT GTG TCA AAG TCA TCT TAT CCC	6401
Gln Gln Ala Met Trp Met Met Thr Ala Val Ser Lys Ser Ser Tyr Pro	
2010 2015 2020	
ATG CGT GTG AAC AGA TGC AAG GAA ATC CTC AAT AAA GCT ATT CAT ATG	6449
Met Arg Val Asn Arg Cys Lys Glu Ile Leu Asn Lys Ala Ile His Met	
2025 2030 2035	
AAA AAA TCC TTA GAG AAG TTT GTT GGA GAT GCA ACT CGC CTA ACA GAT	6497
Lys Lys Ser Leu Glu Lys Phe Val Gly Asp Ala Thr Arg Leu Thr Asp	
2040 2045 2050 2055	
AAG CTT CTA GAA TTG TGC AAT AAA CCG GTG GAA ATT CTT GCT TCT CTT	6545
Lys Leu Leu Glu Leu Cys Asn Lys Pro Val Glu Ile Leu Ala Ser Leu	
2060 2065 2070	
CAG AAA CCA AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG TTC TAC	6593
Gln Lys Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr	
2075 2080 2085	
ATC ATG ATG TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT AGA CTA	6641
Ile Met Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu	
2090 2095 2100	
ATG GAA TTC AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT GCA GAG	6689
Met Glu Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu	
2105 2110 2115	

-60-

TCT CGT AGA AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT CCA CTA Ser Arg Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu 2120 2125 2130 2135	6737
AAT GAT GAA TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT GGT TTG Asn Asp Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu 2140 2145 2150	6785
AGA CCT ATT CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT ATG ACA Arg Pro Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr 2155 2160 2165	6833
GGA AAA GAA CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT TTA TCT Gly Lys Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser 2170 2175 2180	6881
GAA AAA CTC AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT CCT CCT Glu Lys Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro 2185 2190 2195	6929
ATT TTT CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA TCA TGG Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp 2200 2205 2210 2215	6977
TAC AGT AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA ATG Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met 2220 2225 2230	7025
GTT GGT TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT CTC Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu 2235 2240 2245	7073
TTT GAT TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT CTT Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu 2250 2255 2260	7121
TTC AAT AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT CGC Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg 2265 2270 2275	7169
CTG ACT CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG GGT Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly 2280 2285 2290 2295	7217
CTT TTT CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT CAG Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln 2300 2305 2310	7265
CGA GAG CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT CTT Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu 2315 2320 2325	7313
GTG GAA TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG AAT Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn 2330 2335 2340	7361
GAA ACT GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT GAC Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp 2345 2350 2355	7409
ATT GAG CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG ACA Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr 2360 2365 2370 2375	7457
GGA CTG CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA GAA Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu 2380 2385 2390	7505

-61-

GCT ACT GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT CCA 7553  
 Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro  
 2395 2400 2405

TAT ATG TGAAATGAAA TTATGTAAAA GAATATGTTA ATAATCTAAA AGTAAAAAAA 7609  
 Tyr Met

AAAAAAAAAA AA 7621

## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2409 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser  
 1 5 10 15

Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu  
 20 25 30

Met Ser Met Gln Lys Leu Glu Phe Ile Glu Val Thr Leu Leu Thr Val  
 35 40 45

Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu  
 50 55 60

Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile  
 65 70 75 80

Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly  
 85 90 95

Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu  
 100 105 110

Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu  
 115 120 125

Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu  
 130 135 140

Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys  
 145 150 155 160

Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp  
 165 170 175

Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe  
 180 185 190

Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val  
 195 200 205

Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly Ile Glu Val Asp  
 210 215 220

Ala Glu Tyr Leu Leu Gly Pro Leu Tyr Ala Ala Leu Lys Met Glu Ser  
 225 230 235 240

-62-

Met Glu Ile Ile Glu Glu Ile Gln Cys Gln Thr Gln Gln Glu Asn Leu  
 245 250 255  
 Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg Arg Leu Ser Ser  
 260 265 270  
 Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr Glu Glu Ile Lys  
 275 280 285  
 His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser Ala Leu Lys Gln  
 290 295 300  
 Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser Gly Leu Lys Asn  
 305 310 315 320  
 Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val Leu Gln Leu Thr  
 325 330 335  
 Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met Asn Cys Arg Thr  
 340 345 350  
 Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro Ser Val Val Ile  
 355 360 365  
 Thr Trp Met Ser Leu Asp Phe Tyr Thr Thr Val Leu Lys Ser Cys Arg  
 370 375 380  
 Arg Leu Leu Glu Ser Val Gln Lys Arg Thr Gly Gly Asn Ile Asp Lys  
 385 390 395 400  
 Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met Gln Val Asn Ser Ser  
 405 410 415  
 Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly Met Leu Ser Leu Pro  
 420 425 430  
 Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu Lys Leu Thr Thr Phe  
 435 440 445  
 Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile Ser Asp Ser Tyr Ser  
 450 455 460  
 Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu Thr Leu Phe Pro Arg  
 465 470 475 480  
 Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr Asn Trp Ala Leu Gln  
 485 490 495  
 Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe Ile  
 500 505 510  
 Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu Ile  
 515 520 525  
 Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala Ser  
 530 535 540  
 Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu Thr  
 545 550 555 560  
 Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu Phe  
 565 570 575  
 Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser Gln  
 580 585 590

Leu	Lys	Ala	Ser	Val	Cys	Lys	Pro	Phe	Leu	Phe	Leu	Lys	Lys	Lys
		595					600					605		
Ile	Pro	Ser	Pro	Val	Lys	Leu	Ala	Phe	Ile	Asp	Asn	Leu	His	His
	610					615					620			Leu
Cys	Lys	His	Leu	Asp	Phe	Arg	Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala
625					630					635				Val
Leu	Gly	Thr	Leu	Leu	Asn	Leu	Met	Glu	Asp	Pro	Asp	Lys	Asp	Val
				645					650					655
Val	Ala	Phe	Ser	Gly	Asn	Ile	Lys	His	Ile	Leu	Glu	Ser	Leu	Asp
			660					665					670	Ser
Glu	Asp	Gly	Phe	Ile	Lys	Glu	Leu	Phe	Val	Leu	Arg	Met	Lys	Glu
		675					680					685		Ala
Tyr	Thr	His	Ala	Gln	Ile	Ser	Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr
	690					695					700			Leu
Ile	Leu	Thr	Thr	Gly	Asp	Ile	Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu
705					710					715				Val
Pro	Phe	Ala	Leu	Leu	His	Leu	Leu	His	Cys	Leu	Leu	Ser	Lys	Ser
				725					730					735
Ser	Val	Ser	Gly	Ala	Ala	Tyr	Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala
			740					745					750	Ala
Lys	Ser	Val	Lys	Leu	Gln	Ser	Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro
		755					760					765		Ile
Cys	Gln	Phe	Leu	Val	Glu	Ser	Leu	His	Ser	Ser	Gln	Met	Thr	Ala
	770					775					780			Leu
Pro	Asn	Thr	Pro	Cys	Gln	Asn	Ala	Asp	Val	Arg	Lys	Gln	Asp	Val
785					790					795				800
His	Gln	Arg	Glu	Met	Ala	Leu	Asn	Thr	Leu	Ser	Glu	Ile	Ala	Asn
				805					810					815
Phe	Asp	Phe	Pro	Asp	Leu	Asn	Arg	Phe	Leu	Thr	Arg	Thr	Leu	Gln
			820					825					830	Val
Leu	Leu	Pro	Asp	Leu	Ala	Ala	Lys	Ala	Ser	Pro	Ala	Ala	Ser	Ala
		835					840					845		Leu
Ile	Arg	Thr	Leu	Gly	Lys	Gln	Leu	Asn	Val	Asn	Arg	Arg	Glu	Ile
	850					855					860			Leu
Ile	Asn	Asn	Phe	Lys	Tyr	Ile	Phe	Ser	His	Leu	Val	Cys	Ser	Cys
865					870					875				880
Lys	Asp	Glu	Leu	Glu	Arg	Ala	Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr
				885					890					895
Ile	Glu	Leu	Gly	Ser	Leu	Leu	Arg	Gln	Asp	Phe	Gln	Gly	Leu	His
			900					905					910	Asn
Glu	Leu	Leu	Leu	Arg	Ile	Gly	Glu	His	Tyr	Gln	Gln	Val	Phe	Asn
			915				920					925		Gly
Leu	Ser	Ile	Leu	Ala	Ser	Phe	Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln
	930					935					940			Gly

-64-

Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln Pro  
 945 950 955 960  
 Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser Ser  
 965 970 975  
 Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met Ser  
 980 985 990  
 Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val Lys  
 995 1000 1005  
 Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe Pro  
 1010 1015 1020  
 Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp His  
 1025 1030 1035 1040  
 Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu Pro  
 1045 1050 1055  
 Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr Leu  
 1060 1065 1070  
 Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile Tyr  
 1075 1080 1085  
 Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu Gln  
 1090 1095 1100  
 Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr Leu  
 1105 1110 1115 1120  
 Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg Ile  
 1125 1130 1135  
 His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu Lys  
 1140 1145 1150  
 Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile Ser  
 1155 1160 1165  
 Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser Gln  
 1170 1175 1180  
 Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile Asp  
 1185 1190 1195 1200  
 Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp Phe  
 1205 1210 1215  
 Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu Leu  
 1220 1225 1230  
 Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg Ala  
 1235 1240 1245  
 Pro Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr Asp  
 1250 1255 1260  
 Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg Arg  
 1265 1270 1275 1280  
 Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr Arg  
 1285 1290 1295

-65-

Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys Pro  
 1300 1305 1310

Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala Ser  
 1315 1320 1325

Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser Lys  
 1330 1335 1340

Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val Thr  
 1345 1350 1355 1360

Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys Asn  
 1365 1370 1375

Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu Lys  
 1380 1385 1390

His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp Leu  
 1395 1400 1405

Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu Thr  
 1410 1415 1420

Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys Pro  
 1425 1430 1435 1440

His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr Val  
 1445 1450 1455

Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile Pro  
 1460 1465 1470

Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr Arg  
 1475 1480 1485

Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn Ile  
 1490 1495 1500

Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His Glu  
 1505 1510 1515 1520

Pro Asp Gly Val Ser Gly Val Ser Ala Ile Arg Lys Ala Glu Pro Ser  
 1525 1530 1535

Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg Asp  
 1540 1545 1550

Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln Ile  
 1555 1560 1565

Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln Leu  
 1570 1575 1580

Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser Glu  
 1585 1590 1595 1600

Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys Leu  
 1605 1610 1615

Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys Ser  
 1620 1625 1630

Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys Lys  
 1635 1640 1645

-66-

Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala Glu  
 1650 1655 1660

Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr Gln  
 1665 1670 1675 1680

Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu Glu  
 1685 1690 1695

His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser Gln  
 1700 1705 1710

Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn Ser  
 1715 1720 1725

Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu Ser  
 1730 1735 1740

Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp Leu  
 1745 1750 1755 1760

Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala Tyr  
 1765 1770 1775

Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr Val  
 1780 1785 1790

Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala Leu  
 1795 1800 1805

Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu Thr  
 1810 1815 1820

Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu Leu  
 1825 1830 1835 1840

Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala Ile  
 1845 1850 1855

Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu Asp  
 1860 1865 1870

Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met Val  
 1875 1880 1885

Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile Val  
 1890 1895 1900

Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr Gln  
 1905 1910 1915 1920

Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys Ala  
 1925 1930 1935

Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg Asn  
 1940 1945 1950

Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr Leu  
 1955 1960 1965

Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg Ile  
 1970 1975 1980

Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn Ser  
 1985 1990 1995 2000



-67-

Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr Ala  
 2005 2010 2015  
 Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu Ile  
 2020 2025 2030  
 Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val Gly  
 2035 2040 2045  
 Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys Pro  
 2050 2055 2060  
 Val Glu Ile Leu Ala Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu Lys  
 2065 2070 2075 2080  
 Gly Ser Asp Gly Lys Phe Tyr Ile Met Met Cys Lys Pro Lys Asp Asp  
 2085 2090 2095  
 Leu Arg Lys Asp Cys Arg Leu Met Glu Phe Asn Ser Leu Ile Asn Lys  
 2100 2105 2110  
 Cys Leu Arg Lys Asp Ala Glu Ser Arg Arg Arg Glu Leu His Ile Arg  
 2115 2120 2125  
 Thr Tyr Ala Val Ile Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu Trp  
 2130 2135 2140  
 Val Asn Asn Thr Ala Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr Lys  
 2145 2150 2155 2160  
 Glu Lys Gly Val Tyr Met Thr Gly Lys Glu Leu Arg Gln Cys Met Leu  
 2165 2170 2175  
 Pro Lys Ser Ala Ala Leu Ser Glu Lys Leu Lys Val Phe Arg Glu Phe  
 2180 2185 2190  
 Leu Leu Pro Arg His Pro Pro Ile Phe His Glu Trp Phe Leu Arg Thr  
 2195 2200 2205  
 Phe Pro Asp Pro Thr Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg  
 2210 2215 2220  
 Ser Thr Ala Val Met Ser Met Val Gly Tyr Ile Leu Gly Leu Gly Asp  
 2225 2230 2235 2240  
 Arg His Gly Glu Asn Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys Val  
 2245 2250 2255  
 His Val Asp Phe Asn Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu Val  
 2260 2265 2270  
 Pro Glu Ile Val Pro Phe Arg Leu Thr His Asn Met Val Asn Gly Met  
 2275 2280 2285  
 Gly Pro Met Gly Thr Glu Gly Leu Phe Arg Arg Ala Cys Glu Val Thr  
 2290 2295 2300  
 Met Arg Leu Met Arg Asp Gln Arg Glu Pro Leu Met Ser Val Leu Lys  
 2305 2310 2315 2320  
 Thr Phe Leu His Asp Pro Leu Val Glu Trp Ser Lys Pro Val Lys Gly  
 2325 2330 2335  
 His Ser Lys Ala Pro Leu Asn Glu Thr Gly Glu Val Val Asn Glu Lys  
 2340 2345 2350

-68-

Ala Lys Thr His Val Leu Asp Ile Glu Gln Arg Leu Gln Gly Val Ile  
 2355 2360 2365

Lys Thr Arg Asn Arg Val Thr Gly Leu Pro Leu Ser Ile Glu Gly His  
 2370 2375 2380

Val His Tyr Leu Ile Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln  
 2385 2390 2395 2400

Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2405

## (2) INFORMATION FOR SEQ ID NO:3:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2835 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: 517

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..2610

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GTG GAA GCA GCT TGG AAA TTG TCA CAG TGG GAT TTG GTG GAA AAC TAT	48
Val Glu Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr	
1 5 10 15	
TTG GCA GCA GAT GGA AAA TCT ACA ACA TGG AGT GTC AGA CTG GGA CAG	96
Leu Ala Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg Leu Gly Gln	
20 25 30	
CTA TTA TTA TCA GCC AAA AAA AGA GAT ATC ACA GCT TTT TAT GAC TCA	144
Leu Leu Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser	
35 40 45	
CTG AAA CTA GTG AGA GCA GAA CAA ATT GTA CCT CTT TCA GCT GCA AGC	192
Leu Lys Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser Ala Ala Ser	
50 55 60	
TTT GAA AGA GGC TCC TAC CAA CGA GGA TAT GAA TAT ATT GTG AGA TTG	240
Phe Glu Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu	
65 70 75 80	
CAC ATG TTA TGT GAG TTG GAG CAT AGC ATC AAA CCA CTT TTC CAG CAT	288
His Met Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu Phe Gln His	
85 90 95	
TCT CCA GGT GAC AGT TCT CAA GAA GAT TCT CTA AAC TGG GTA GCT CGA	336
Ser Pro Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp Val Ala Arg	
100 105 110	
CTA GAA ATG ACC CAG AAT TCC TAC AGA GCC AAG GAG CCT ATC CTG GCT	384
Leu Glu Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala	
115 120 125	

-69-

CTC Leu 130	CGG Arg	AGG Arg	GCT Ala	TTA Leu	CTA Leu	AGC Ser 135	CTC Leu	AAC Asn	AAA Lys	AGA Arg	CCA Pro 140	GAT Asp	TAC Tyr	AAT Asn	GAA Glu	432
ATG Met 145	GTT Val	GGA Gly	GAA Glu	TGC Cys	TGG Trp 150	CTG Leu	CAG Gln	AGT Ser	GCC Ala	AGG Arg 155	GTA Val	GCT Ala	AGA Arg	AAG Lys	GCT Ala 160	480
GGT Gly	CAC His	CAC His	CAG Gln	ACA Thr 165	GCC Ala	TAC Tyr	AAT Asn	GCT Ala	CTC Leu 170	CTT Leu	AAT Asn	GCA Ala	GGG Gly	GAA Glu 175	TCA Ser	528
CGA Arg	CTC Leu	GCT Ala	GAA Glu 180	CTG Leu	TAC Tyr	GTG Val	GAA Glu 185	AGG Arg	GCA Ala	AAG Lys	TGG Trp	CTC Leu	TGG Trp 190	TCC Ser	AAG Lys	576
GGT Gly	GAT Asp	GTT Val 195	CAC His	CAG Gln	GCA Ala	CTA Leu 200	ATT Ile	GTT Val	CTT Leu	CAA Gln	AAA Lys 205	GGT Gly 205	GTT Val	GAA Glu	TTA Leu	624
TGT Cys	TTT Phe 210	CCT Pro	GAA Glu	AAT Asn	GAA Glu	ACC Thr 215	CCA Pro	CCT Pro	GAG Glu	GGT Gly	AAG Lys 220	AAC Asn	ATG Met	TTA Leu	ATC Ile	672
CAT His 225	GGT Gly	CGA Arg	GCT Ala	ATG Met	CTA Leu 230	CTA Leu	GTG Val	GGC Gly	CGA Arg	TTT Phe 235	ATG Met	GAA Glu	GAA Glu	ACA Thr	GCT Ala 240	720
AAC Asn	TTT Phe	GAA Glu	AGC Ser	AAT Asn 245	GCA Ala	ATT Ile	ATG Met	AAA Lys	AAA Lys 250	TAT Tyr	AAG Lys	GAT Asp	GTG Val	ACC Thr 255	GCG Ala	768
TGC Cys	CTG Leu	CCA Pro	GAA Glu 260	TGG Trp	GAG Glu	GAT Asp	GGG Gly	CAT His 265	TTT Phe	TAC Tyr	CTT Leu	GCC Ala	AAG Lys 270	TAC Tyr	TAT Tyr	816
GAC Asp	AAA Lys	TTG Leu 275	ATG Met	CCC Pro	ATG Met	GTC Val	ACA Thr 280	GAC Asp	AAC Asn	AAA Lys	ATG Met	GAA Glu 285	AAG Lys	CAA Gln	GGT Gly	864
GAT Asp	CTC Leu	ATC Ile	CGG Arg	TAT Tyr	ATA Ile	GTT Val 295	CTT Leu	CAT His	TTT Phe	GGC Gly	AGA Arg 300	TCT Ser	CTA Leu	CAA Gln	TAT Tyr	912
GGA Gly 305	AAT Asn	CAG Gln	TTC Phe	ATA Ile	TAT Tyr 310	CAG Gln	TCA Ser	ATG Met	CCA Pro	CGA Arg 315	ATG Met	TTA Leu	ACT Thr	CTA Leu	TGG Trp 320	960
CTT Leu	GAT Asp	TAT Tyr	GGT Gly	ACA Thr 325	AAG Lys	TCA Ser	TAT Tyr	GAA Glu	TGG Trp 330	GAA Glu	AAA Lys	GCT Ala	GGC Gly	CGC Arg 335	TCC Ser	1008
GAT Asp	CGT Arg	GTA Val	CAA Gln 340	ATG Met	AGG Arg	AAT Asn	GAT Asp	TTG Leu 345	GGT Gly	AAA Lys	ATA Ile	AAC Asn	AAG Lys 350	GTT Val	ATC Ile	1056
ACA Thr	GAG Glu	CAT His 355	ACA Thr	AAC Asn	TAT Tyr	TTA Leu	GCT Ala 360	CCA Pro	TAT Tyr	CAA Gln	TTT Phe 365	TTG Leu	ACT Thr	GCT Ala	TTT Phe	1104
TCA Ser	CAA Gln 370	TTG Leu	ATC Ile	TCT Ser	CGA Arg	ATT Ile 375	TGT Cys	CAT His	TCT Ser	CAC His	GAT Asp 380	GAA Glu	GTT Val	TTT Phe	GTT Val	1152

-70-

GTC Val 385	TTG Leu	ATG Met	GAA Glu	ATA Ile	ATA Ile 390	GCC Ala	AAA Lys	GTA Val	TTT Phe 395	CTA Leu	GCC Ala	TAT Tyr	CCT Pro	CAA Gln 400	CAA Gln	1200
GCA Ala	ATG Met	TGG Trp	ATG Met	ATG Met 405	ACA Thr	GCT Ala	GTG Val	TCA Ser	AAG Lys 410	TCA Ser	TCT Ser	TAT Tyr	CCC Pro	ATG Met 415	CGT Arg	1248
GTG Val	AAC Asn	AGA Arg	TGC Cys 420	AAG Lys	GAA Glu	ATC Ile	CTC Leu	AAT Asn 425	AAA Lys	GCT Ala	ATT Ile	CAT His	ATG Met 430	AAA Lys	AAA Lys	1296
TCC Ser	TTA Leu	GAG Glu 435	AAG Lys	TTT Phe	GTT Val	GGA Gly	GAT Asp 440	GCA Ala	ACT Thr	CGC Arg	CTA Leu	ACA Thr 445	GAT Asp	AAG Lys	CTT Leu	1344
CTA Leu 450	GAA Glu	TTG Leu	TGC Cys	AAT Asn	AAA Lys	CCG Pro 455	GTT Val	GAT Asp	GGA Gly	AGT Ser	AGT Ser	TCC Ser	ACA Thr	TTA Leu	AGC Ser	1392
ATG Met 465	AGC Ser	ACT Thr	CAT His	TTT Phe	AAA Lys 470	ATG Met	CTT Leu	AAA Lys	AAG Lys	CTG Leu 475	GTA Val	GAA Glu	GAA Glu	GCA Ala	ACA Thr 480	1440
TTT Phe	AGT Ser	GAA Glu	ATC Ile	CTC Leu 485	ATT Ile	CCT Pro	CTA Leu	CAA Gln 490	TCA Ser	GTC Val	ATG Met	ATA Ile	CCT Pro	ACA Thr 495	CTT Leu	1488
CCA Pro	TCA Ser	ATT Ile 500	CTG Leu	GGT Gly	ACC Thr	CAT His	GCT Ala	AAC Asn 505	CAT His	GCT Ala	AGC Ser	CAT His	GAA Glu 510	CCA Pro	TTT Phe	1536
CCT Pro	GGA Gly	CAT His 515	TGG Trp	GCC Ala	TAT Tyr	ATT Ile	GCA Ala 520	GGG Gly	TTT Phe	GAT Asp	GAT Asp	ATG Met 525	GTG Val	GAA Glu	ATT Ile	1584
CTT Leu 530	GCT Ala	TCT Ser	CTT Leu	CAG Gln	AAA Lys	CCA Pro 535	AAG Lys	AAG Lys	ATT Ile	TCT Ser	TTA Leu 540	AAA Lys	GGC Gly	TCA Ser	GAT Asp	1632
GGA Gly 545	AAG Lys	TTC Phe	TAC Tyr	ATC Ile	ATG Met 550	ATG Met	TGT Cys	AAG Lys	CCA Pro	AAA Lys 555	GAT Asp	GAC Asp	CTG Leu	AGA Arg	AAG Lys 560	1680
GAT Asp	TGT Cys	AGA Arg	CTA Leu	ATG Met 565	GAA Glu	TTC Phe	AAT Asn	TCC Ser	TTG Leu 570	ATT Ile	AAT Asn	AAG Lys	TGC Cys	TTA Leu 575	AGA Arg	1728
AAA Lys	GAT Asp	GCA Ala	GAG Glu	TCT Ser	CGT Arg	AGA Arg	AGA Arg	GAA Glu 585	CTT Leu	CAT His	ATT Ile	CGA Arg	ACA Thr 590	TAT Tyr	GCA Ala	1776
GTT Val	ATT Ile	CCA Pro 595	CTA Leu	AAT Asn	GAT Asp	GAA Glu	TGT Cys 600	GGG Gly	ATT Ile	ATT Ile	GAA Glu	TGG Trp 605	GTG Val	AAC Asn	AAC Asn	1824
ACT Thr 610	GCT Ala	GGT Gly	TTG Leu	AGA Arg	CCT Pro	ATT Ile 615	CTG Leu	ACC Thr	AAA Lys	CTA Leu	TAT Tyr 620	AAA Lys	GAA Glu	AAG Lys	GGA Gly	1872
GTG Val 625	TAT Tyr	ATG Met	ACA Thr	GGA Gly	AAA Lys 630	GAA Glu	CTT Leu	CGC Arg	CAG Gln	TGT Cys 635	ATG Met	CTA Leu	CCA Pro	AAG Lys	TCA Ser 640	1920
GCA Ala	GCT Ala	TTA Leu	TCT Ser	GAA Glu 645	AAA Lys	CTC Leu	AAA Lys	GTA Val	TTC Phe 650	CGA Arg	GAA Glu	TTT Phe	CTC Leu	CTG Leu 655	CCC Pro	1968

-71-

AGG CAT CCT CCT ATT TTT CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT	2016
Arg His Pro Pro Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp	
660 665 670	
CCT ACA TCA TGG TAC AGT AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA	2064
Pro Thr Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala	
675 680 685	
GTA ATG TCA ATG GTT GGT TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT	2112
Val Met Ser Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly	
690 695 700	
GAA AAT ATT CTC TTT GAT TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT	2160
Glu Asn Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp	
705 710 715 720	
TTC AAT TGT CTT TTC AAT AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT	2208
Phe Asn Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile	
725 730 735	
GTG CCA TTT CGC CTG ACT CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG	2256
Val Pro Phe Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met	
740 745 750	
GGA ACA GAG GGT CTT TTT CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG	2304
Gly Thr Glu Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu	
755 760 765	
ATG CGT GAT CAG CGA GAG CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA	2352
Met Arg Asp Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu	
770 775 780	
CAT GAT CCT CTT GTG GAA TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA	2400
His Asp Pro Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys	
785 790 795 800	
GCG CCA CTG AAT GAA ACT GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC	2448
Ala Pro Leu Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr	
805 810 815	
CAT GTT CTT GAC ATT GAG CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA	2496
His Val Leu Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg	
820 825 830	
AAT AGA GTG ACA GGA CTG CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC	2544
Asn Arg Val Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr	
835 840 845	
CTT ATA CAA GAA GCT ACT GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT	2592
Leu Ile Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu	
850 855 860	
GGT TGG ACT CCA TAT ATG TGAAATGAAA TTATGTAAAA GAATATGTTA	2640
Gly Trp Thr Pro Tyr Met	
865 870	
ATAATCTAAA AGTAATGCAT TTGGTATGAA TCTGTGGTTG TATCTGTTCA ATTCTAAAGT	2700
ACAACATAAA TTTACGTTCT CAGCAACTGT TATTTCTCTC TGATCATTAA TTATATGTAA	2760
AATAATATAC ATTCAATTAT TAAGAAATAA ACTGCTTTCT TAATAAAAAA AAAAAAAA	2820
AAAAAAAAA AAAAA	2835

-72-

## (2) INFORMATION FOR SEQ ID NO:4:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 870 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

```

Val Glu Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr
  1          5          10          15
Leu Ala Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg Leu Gly Gln
          20          25          30
Leu Leu Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser
          35          40          45
Leu Lys Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser Ala Ala Ser
          50          55          60
Phe Glu Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu
          65          70          75          80
His Met Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu Phe Gln His
          85          90          95
Ser Pro Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp Val Ala Arg
          100          105          110
Leu Glu Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala
          115          120          125
Leu Arg Arg Ala Leu Leu Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu
          130          135          140
Met Val Gly Glu Cys Trp Leu Gln Ser Ala Arg Val Ala Arg Lys Ala
          145          150          155          160
Gly His His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser
          165          170          175
Arg Leu Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu Trp Ser Lys
          180          185          190
Gly Asp Val His Gln Ala Leu Ile Val Leu Gln Lys Gly Val Glu Leu
          195          200          205
Cys Phe Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn Met Leu Ile
          210          215          220
His Gly Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu Glu Thr Ala
          225          230          235          240
Asn Phe Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp Val Thr Ala
          245          250          255
Cys Leu Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr
          260          265          270
Asp Lys Leu Met Pro Met Val Thr Asp Asn Lys Met Glu Lys Gln Gly
          275          280          285

```

-73-

Asp Leu Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser Leu Gln Tyr  
 290 295 300  
 Gly Asn Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu Thr Leu Trp  
 305 310 315 320  
 Leu Asp Tyr Gly Thr Lys Ser Tyr Glu Trp Glu Lys Ala Gly Arg Ser  
 325 330 335  
 Asp Arg Val Gln Met Arg Asn Asp Leu Gly Lys Ile Asn Lys Val Ile  
 340 345 350  
 Thr Glu His Thr Asn Tyr Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe  
 355 360 365  
 Ser Gln Leu Ile Ser Arg Ile Cys His Ser His Asp Glu Val Phe Val  
 370 375 380  
 Val Leu Met Glu Ile Ile Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln  
 385 390 395 400  
 Ala Met Trp Met Met Thr Ala Val Ser Lys Ser Ser Tyr Pro Met Arg  
 405 410 415  
 Val Asn Arg Cys Lys Glu Ile Leu Asn Lys Ala Ile His Met Lys Lys  
 420 425 430  
 Ser Leu Glu Lys Phe Val Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu  
 435 440 445  
 Leu Glu Leu Cys Asn Lys Pro Val Asp Gly Ser Ser Ser Thr Leu Ser  
 450 455 460  
 Met Ser Thr His Phe Lys Met Leu Lys Lys Leu Val Glu Glu Ala Thr  
 465 470 475 480  
 Phe Ser Glu Ile Leu Ile Pro Leu Gln Ser Val Met Ile Pro Thr Leu  
 485 490 495  
 Pro Ser Ile Leu Gly Thr His Ala Asn His Ala Ser His Glu Pro Phe  
 500 505 510  
 Pro Gly His Trp Ala Tyr Ile Ala Gly Phe Asp Asp Met Val Glu Ile  
 515 520 525  
 Leu Ala Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp  
 530 535 540  
 Gly Lys Phe Tyr Ile Met Met Cys Lys Pro Lys Asp Asp Leu Arg Lys  
 545 550 555 560  
 Asp Cys Arg Leu Met Glu Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg  
 565 570 575  
 Lys Asp Ala Glu Ser Arg Arg Arg Glu Leu His Ile Arg Thr Tyr Ala  
 580 585 590  
 Val Ile Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu Trp Val Asn Asn  
 595 600 605  
 Thr Ala Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly  
 610 615 620  
 Val Tyr Met Thr Gly Lys Glu Leu Arg Gln Cys Met Leu Pro Lys Ser  
 625 630 635 640

-74-

Ala Ala Leu Ser Glu Lys Leu Lys Val Phe Arg Glu Phe Leu Leu Pro  
645 650 655

Arg His Pro Pro Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp  
660 665 670

Pro Thr Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala  
675 680 685

Val Met Ser Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly  
690 695 700

Glu Asn Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp  
705 710 715 720

Phe Asn Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile  
725 730 735

Val Pro Phe Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met  
740 745 750

Gly Thr Glu Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu  
755 760 765

Met Arg Asp Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu  
770 775 780

His Asp Pro Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys  
785 790 795 800

Ala Pro Leu Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr  
805 810 815

His Val Leu Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg  
820 825 830

Asn Arg Val Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr  
835 840 845

Leu Ile Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu  
850 855 860

Gly Trp Thr Pro Tyr Met  
865 870

## (2) INFORMATION FOR SEQ ID NO:5:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 33 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: Primer oDH15a

## (ix) FEATURE:

- (A) NAME/KEY: modified\_base
- (B) LOCATION: group(15, 18, 24, 30)
- (D) OTHER INFORMATION: The nucleotides at these positions are inosines.



-75-

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

GCAGACGGAT CCGGNWCNGA YGGNAAYHTN TAY

33

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 27 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(ix) FEATURE:

- (A) NAME/KEY: modified\_base
- (B) LOCATION: group(15, 18, 24)
- (D) OTHER INFORMATION: The nucleotides at these positions are inosines.

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

GCAGACGGAT CCGGNWCNGA YGGNAAY

27

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 30 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(vii) IMMEDIATE SOURCE:

- (B) CLONE: Primer oDH16

(ix) FEATURE:

- (A) NAME/KEY: modified\_base
- (B) LOCATION: 24
- (D) OTHER INFORMATION: The nucleotides at these positions are inosines.

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

GCAGACGAAT TCRCARTYRA ARTCNACRTG

30

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 41 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(vii) IMMEDIATE SOURCE:

- (B) CLONE: Primer oDH17a

-76-

(ix) FEATURE:  
    (A) NAME/KEY: modified\_base  
    (B) LOCATION: group(21, 24, 27, 30)  
    (D) OTHER INFORMATION: The nucleotides at these positions are inosines.

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:  
GCAGACGGAT CCAARTTYCC NCCNRTNYTN TAYSARTGGT T 41

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 41 base pairs  
    (B) TYPE: nucleic acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(vii) IMMEDIATE SOURCE:  
    (B) CLONE: Primer oDH17b

(ix) FEATURE:  
    (A) NAME/KEY: modified\_base  
    (B) LOCATION: group(24, 27, 30, 33)  
    (D) OTHER INFORMATION: The nucleotides at these positions are inosines.

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:  
GCAGACGAAT CCAACCAYS RTANARNAYN GGNGGAAAYT T 41

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 32 base pairs  
    (B) TYPE: nucleic acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(vii) IMMEDIATE SOURCE:  
    (B) CLONE: Primer oDH18a

(ix) FEATURE:  
    (A) NAME/KEY: modified\_base  
    (B) LOCATION: group(15, 18, 21, 24, 30)  
    (D) OTHER INFORMATION: The nucleotides at these positions are inosines.

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:  
GCAGACGGAT CCYTNGGNYT NGNGAYCGN CA 32

-77-

## (2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 32 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer oDH18b

- (ix) FEATURE:
  - (A) NAME/KEY: modified\_base
  - (B) LOCATION: group(15, 18, 21)
  - (D) OTHER INFORMATION: The nucleotides at these positions are

inosines.

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

GCAGACGGAT CCYTNGGNYT NGGNGAYAGR CA

32

## (2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 33 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer oDH23

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

GACGCAGAAT TCACCAAGTCA AAGAATCAAA GAG

33

## (2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 16 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer mo3

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

CTACAGAGCC AAGGAG

16

-78-

## (2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 22 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer mo6

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

TCGAGCTATG CTACTAGTGG GC

22

## (2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 17 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer oHT9-1

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

CCAGTAAACT TGCTTTC

17

## (2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 20 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: Primer oHT9-4

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

TTTGC GGCCC TTCCAATATC

20

## (2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 7440 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

-79-

(vii) IMMEDIATE SOURCE:  
(B) CLONE: MCCS1beta

(ix) FEATURE:  
(A) NAME/KEY: CDS  
(B) LOCATION: 1..7437

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

ATG GGT CAT GCT GTG GAA TGG CCA GTG GTC ATG AGC CGA TTT TTA AGT	48
Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser	
1 5 10 15	
CAA TTA GAT GAA CAC ATG GGA TAT TTA CAA TCA GCT CCT TTG CAG TTG	96
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu	
20 25 30	
ATG AGT ATG CAA AAA TTA GAA TTT ATT GAA GTC ACT TTA TTA ACG GTT	144
Met Ser Met Gln Lys Leu Glu Phe Ile Glu Val Thr Leu Leu Thr Val	
35 40 45	
CTT ACT CGT ATT ATT GCA ATT GTG TTT TTT AGA AGG CAA GAA CTC TTA	192
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu	
50 55 60	
CTT TGG CAG ATA GGT TGT GTT CTG CTA GAG TAT GGT AGT CCA AAA ATT	240
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile	
65 70 75 80	
AAA TCC CTA GCA ATT AGC TTT TTA ACA GAA CTT TTT CAG CTT GGA GGA	288
Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly	
85 90 95	
CTA CCA GCA CAA CCA GCT AGC ACT TTT TTC AGC TCA TTT TTG GAA TTA	336
Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu	
100 105 110	
TTA AAA CAC CTT GTA GAA ATG GAT ACT GAC CAA TTG AAA CTC TAT GAA	384
Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu	
115 120 125	
GAG CCA TTA TCA AAG CTG ATA AAG ACA CTA TTT CCC TTT GAA GCA GAA	432
Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu	
130 135 140	
GCT TAT AGA AAT ATT GAA CCT GTC TAT TTA AAT ATG CTG CTG GAA AAA	480
Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys	
145 150 155 160	
CTC TGT GTC ATG TTT GAA GAC GGT GTG CTC ATG CGG CTT AAG TCT GAT	528
Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp	
165 170 175	
TTG CTA AAA GCA GCT TTG TGC CAT TTA CTG CAG TAT TTC CTT AAA TTT	576
Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe	
180 185 190	
GTG CCA GCT GGG TAT GAA TCT GCT TTA CAA GTC AGG AAG GTC TAT GTG	624
Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val	
195 200 205	
AGA AAT ATT TGT AAA GCT CTT TTG GAT GTG CTT GGA ATT GAG GTA GAT	672
Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly Ile Glu Val Asp	
210 215 220	

-80-

GCA Ala 225	GAG Glu 225	TAC Tyr 225	TTG Leu 225	TTG Leu 230	GGC Gly 230	CCA Pro 230	CTT Leu 230	TAT Tyr 230	GCA Ala 235	GCT Ala 235	TTG Leu 235	AAA Lys 235	ATG Met 240	GAA Glu 240	AGT Ser 240	720
ATG Met 245	GAA Glu 245	ATC Ile 245	ATT Ile 245	GAG Glu 245	GAG Glu 245	ATT Ile 245	CAA Gln 250	TGC Cys 250	CAA Gln 250	ACT Thr 250	CAA Gln 255	CAG Gln 255	GAA Glu 255	AAC Asn 255	CTC Leu 255	768
AGC Ser 260	AGT Ser 260	AAT Asn 260	AGT Ser 260	GAT Asp 260	GGA Gly 265	ATA Ile 265	TCA Ser 265	CCC Pro 265	AAA Lys 265	AGG Arg 265	CGT Arg 270	CGT Arg 270	CTC Leu 270	AGC Ser 270	TCG Ser 270	816
TCT Ser 275	CTA Leu 275	AAC Asn 275	CCT Pro 275	TCT Ser 275	AAA Lys 280	AGA Arg 280	GCA Ala 280	CCA Pro 280	AAA Lys 285	CAG Gln 285	ACT Thr 285	GAG Glu 285	GAA Glu 285	ATT Ile 285	AAA Lys 285	864
CAT His 290	GTG Val 290	GAC Asp 290	ATG Met 290	AAC Asn 295	CAA Gln 295	AAG Lys 295	AGC Ser 295	ATA Ile 295	TTA Leu 300	TGG Trp 300	AGT Ser 300	GCA Ala 300	CTG Leu 300	AAA Lys 300	CAG Gln 300	912
AAA Lys 305	GCT Ala 305	GAA Glu 305	TCC Ser 305	CTT Leu 310	CAG Gln 310	ATT Ile 310	TCC Ser 310	CTT Leu 315	GAA Glu 315	TAC Tyr 315	AGT Ser 315	GGC Gly 315	CTA Leu 315	AAG Lys 315	AAT Asn 315	960
CCT Pro 325	GTT Val 325	ATT Ile 325	GAG Glu 325	ATG Met 325	TTA Leu 325	GAA Glu 325	GGA Gly 325	ATT Ile 330	GCT Ala 330	GTT Val 330	GTC Val 330	TTA Leu 330	CAA Gln 330	CTG Leu 330	ACT Thr 330	1008
GCT Ala 340	CTG Leu 340	TGT Cys 340	ACT Thr 340	GTT Val 340	CAT His 345	TGT Cys 345	TCT Ser 345	CAT His 345	CAA Gln 345	AAC Asn 345	ATG Met 345	AAC Asn 345	TGC Cys 350	CGT Arg 350	ACT Thr 350	1056
TTC Phe 355	AAG Lys 355	GAC Asp 355	TGT Cys 355	CAA Gln 360	CAT His 360	AAA Lys 360	TCC Ser 360	AAG Lys 360	AAG Lys 360	AAA Lys 365	CCT Pro 365	TCT Ser 365	GTA Val 365	GTG Val 365	ATA Ile 365	1104
ACT Thr 370	TGG Trp 370	ATG Met 370	TCA Ser 370	TTG Leu 375	GAT Asp 375	TTT Phe 375	TAC Tyr 375	ACA Thr 375	ACA Thr 375	GTG Val 380	CTT Leu 380	AAG Lys 380	AGC Ser 380	TGT Cys 380	AGA Arg 380	1152
AGG Arg 385	TTG Leu 385	TTA Leu 385	GAA Glu 385	TCT Ser 390	GTT Val 390	CAG Gln 390	AAA Lys 390	CGG Arg 395	ACT Thr 395	GGA Gly 395	GGC Gly 395	AAC Asn 395	ATT Ile 395	GAT Asp 400	AAG Lys 400	1200
GTG Val 405	GTG Val 405	AAA Lys 405	ATT Ile 405	TAT Tyr 405	GAT Asp 405	GCT Ala 410	TTG Leu 410	ATT Ile 410	TAT Tyr 410	ATG Met 415	CAA Gln 415	GTA Val 415	AAC Asn 415	AGT Ser 415	TCA Ser 415	1248
TTT Phe 420	GAA Glu 420	GAT Asp 420	CAT His 420	ATC Ile 420	CTG Leu 425	GAA Glu 425	GAT Asp 425	TTA Leu 425	TGT Cys 425	GGA Gly 430	ATG Met 430	CTC Leu 430	TCA Ser 430	CTT Leu 430	CCA Pro 430	1296
TGG Trp 435	ATT Ile 435	TAT Tyr 435	TCC Ser 435	CAT His 435	TCT Ser 440	GAT Asp 440	GAT Asp 440	GGC Gly 440	TGT Cys 440	TTA Leu 445	AAG Lys 445	TTG Leu 445	ACC Thr 445	ACA Thr 445	TTT Phe 445	1344
GCC Ala 450	GCT Ala 450	AAT Asn 450	CTT Leu 450	CTA Leu 455	ACA Thr 455	TTA Leu 455	AGC Ser 455	TGT Cys 455	AGG Arg 460	ATT Ile 460	TCA Ser 460	GAT Asp 460	AGC Ser 460	TAT Tyr 460	TCA Ser 460	1392
CCA Pro 465	CAG Gln 465	GCA Ala 465	CAA Gln 465	TCA Ser 470	CGA Arg 470	TGT Cys 470	GTG Val 470	TTT Phe 475	CTT Leu 475	CTG Leu 475	ACT Thr 475	CTG Leu 475	TTT Phe 475	CCA Pro 480	AGA Arg 480	1440
AGA Arg 485	ATA Ile 485	TTC Phe 485	CTT Leu 485	GAG Glu 485	TGG Trp 485	AGA Arg 485	ACA Thr 485	GCA Ala 490	GTT Val 490	TAC Tyr 490	AAC Asn 490	TGG Trp 490	GCC Ala 495	CTG Leu 495	CAG Gln 495	1488

-81-

AGC	TCC	CAT	GAA	GTA	ATC	CGG	GCT	AGT	TGT	GTT	AGT	GGA	TTT	TTT	ATC	1536
Ser	Ser	His	Glu	Val	Ile	Arg	Ala	Ser	Cys	Val	Ser	Gly	Phe	Phe	Ile	
			500					505					510			
TTA	TTG	CAG	CAG	CAG	AAT	TCT	TGT	AAC	AGA	GTT	CCC	AAG	ATT	CTT	ATA	1584
Leu	Leu	Gln	Gln	Gln	Asn	Ser	Cys	Asn	Arg	Val	Pro	Lys	Ile	Leu	Ile	
		515					520					525				
GAT	AAA	GTC	AAA	GAT	GAT	TCT	GAC	ATT	GTC	AAG	AAA	GAA	TTT	GCT	TCT	1632
Asp	Lys	Val	Lys	Asp	Asp	Ser	Asp	Ile	Val	Lys	Lys	Glu	Phe	Ala	Ser	
	530					535					540					
ATA	CTT	GGT	CAA	CTT	GTC	TGT	ACT	CTT	CAC	GGC	ATG	TTT	TAT	CTG	ACA	1680
Ile	Leu	Gly	Gln	Leu	Val	Cys	Thr	Leu	His	Gly	Met	Phe	Tyr	Leu	Thr	
545					550					555					560	
AGT	TCT	TTA	ACA	GAA	CCT	TTC	TCT	GAA	CAC	GGA	CAT	GTG	GAC	CTC	TTC	1728
Ser	Ser	Leu	Thr	Glu	Pro	Phe	Ser	Glu	His	Gly	His	Val	Asp	Leu	Phe	
				565					570					575		
TGT	AGG	AAC	TTG	AAA	GCC	ACT	TCT	CAA	CAT	GAA	TGT	TCA	TCT	TCT	CAA	1776
Cys	Arg	Asn	Leu	Lys	Ala	Thr	Ser	Gln	His	Glu	Cys	Ser	Ser	Ser	Gln	
			580					585					590			
CTA	AAA	GCT	TCT	GTC	TGC	AAG	CCA	TTC	CTT	TTC	CTA	CTG	AAA	AAA	AAA	1824
Leu	Lys	Ala	Ser	Val	Cys	Lys	Pro	Phe	Leu	Phe	Leu	Leu	Lys	Lys	Lys	
		595					600					605				
ATA	CCT	AGT	CCA	GTA	AAA	CTT	GCT	TTC	ATA	GAT	AAT	CTA	CAT	CAT	CTT	1872
Ile	Pro	Ser	Pro	Val	Lys	Leu	Ala	Phe	Ile	Asp	Asn	Leu	His	His	Leu	
	610					615					620					
TGT	AAG	CAT	CTT	GAT	TTT	AGA	GAA	GAT	GAA	ACA	GAT	GTA	AAA	GCA	GTT	1920
Cys	Lys	His	Leu	Asp	Phe	Arg	Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala	Val	
625					630					635					640	
CTT	GGA	ACT	TTA	TTA	AAT	TTA	ATG	GAA	GAT	CCA	GAC	AAA	GAT	GTT	AGA	1968
Leu	Gly	Thr	Leu	Leu	Asn	Leu	Met	Glu	Asp	Pro	Asp	Lys	Asp	Val	Arg	
				645					650					655		
GTG	GCT	TTT	AGT	GGA	AAT	ATC	AAG	CAC	ATA	TTG	GAA	TCC	TTG	GAC	TCT	2016
Val	Ala	Phe	Ser	Gly	Asn	Ile	Lys	His	Ile	Leu	Glu	Ser	Leu	Asp	Ser	
			660					665					670			
GAA	GAT	GGA	TTT	ATA	AAG	GAG	CTT	TTT	GTC	TTA	AGA	ATG	AAG	GAA	GCA	2064
Glu	Asp	Gly	Phe	Ile	Lys	Glu	Leu	Phe	Val	Leu	Arg	Met	Lys	Glu	Ala	
		675					680					685				
TAT	ACA	CAT	GCC	CAA	ATA	TCA	AGA	AAT	AAT	GAG	CTG	AAG	GAT	ACC	TTG	2112
Tyr	Thr	His	Ala	Gln	Ile	Ser	Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr	Leu	
	690					695					700					
ATT	CTT	ACA	ACA	GGG	GAT	ATT	GGA	AGG	GCC	GCA	AAA	GGA	GAT	TTG	GTA	2160
Ile	Leu	Thr	Thr	Gly	Asp	Ile	Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu	Val	
705					710					715					720	
CCA	TTT	GCA	CTC	TTA	CAC	TTA	TTG	CAT	TGT	TTG	TTA	TCC	AAG	TCA	GCA	2208
Pro	Phe	Ala	Leu	Leu	His	Leu	Leu	His	Cys	Leu	Leu	Ser	Lys	Ser	Ala	
				725					730					735		
TCT	GTC	TCT	GGA	GCA	GCA	TAC	ACA	GAA	ATT	AGA	GCT	CTG	GTT	GCA	GCT	2256
Ser	Val	Ser	Gly	Ala	Ala	Tyr	Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala	Ala	
			740					745					750			
AAA	AGT	GTT	AAA	CTG	CAA	AGT	TTT	TTC	AGC	CAG	TAT	AAG	AAA	CCC	ATC	2304
Lys	Ser	Val	Lys	Leu	Gln	Ser	Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro	Ile	
		755					760					765				

-82-

TGT Cys	CAG Gln	TTT Phe	TTG Leu	GTA Val	GAA Glu	TCC Ser	CTT Leu	CAC His	TCT Ser	AGT Ser	CAG Gln	ATG Met	ACA Thr	GCA Ala	CTT Leu	2352
770						775					780					
CCG Pro	AAT Asn	ACT Thr	CCA Pro	TGC Cys	CAG Gln	AAT Asn	GCT Ala	GAC Asp	GTG Val	CGA Arg	AAA Lys	CAA Gln	GAT Asp	GTG Val	GCT Ala	2400
785					790					795					800	
CAC His	CAG Gln	AGA Arg	GAA Glu	ATG Met	GCT Ala	TTA Leu	AAT Asn	ACG Thr	TTG Leu	TCT Ser	GAA Glu	ATT Ile	GCC Ala	AAC Asn	GTT Val	2448
				805					810					815		
TTC Phe	GAC Asp	TTT Phe	CCT Pro	GAT Asp	CTT Leu	AAT Asn	CGT Arg	TTT Phe	CTT Leu	ACT Thr	AGG Arg	ACA Thr	TTA Leu	CAA Gln	GTT Val	2496
			820					825						830		
CTA Leu	CTA Leu	CCT Pro	GAT Asp	CTT Leu	GCT Ala	GCC Ala	AAA Lys	GCA Ala	AGC Ser	CCT Pro	GCA Ala	GCT Ala	TCT Ser	GCT Ala	CTC Leu	2544
		835					840						845			
ATT Ile	CGA Arg	ACT Thr	TTA Leu	GGA Gly	AAA Lys	CAA Gln	TTA Leu	AAT Asn	GTC Val	AAT Asn	CGT Arg	AGA Arg	GAG Glu	ATT Ile	TTA Leu	2592
	850					855					860					
ATA Ile	AAC Asn	AAC Asn	TTT Phe	AAA Lys	TAT Tyr	ATT Ile	TTT Phe	TCT Ser	CAT His	TTG Leu	GTC Val	TGT Cys	TCT Ser	TGT Cys	TCC Ser	2640
	865				870					875					880	
AAA Lys	GAT Asp	GAA Glu	TTA Leu	GAA Glu	CGT Arg	GCC Ala	CTT Leu	CAT His	TAT Tyr	CTG Leu	AAG Lys	AAT Asn	GAA Glu	ACA Thr	GAA Glu	2688
				885					890					895		
ATT Ile	GAA Glu	CTG Leu	GGG Gly	AGC Ser	CTG Leu	TTG Leu	AGA Arg	CAA Gln	GAT Asp	TTC Phe	CAA Gln	GGA Gly	TTG Leu	CAT His	AAT Asn	2736
			900					905					910			
GAA Glu	TTA Leu	TTG Leu	CTG Leu	CGT Arg	ATT Ile	GGA Gly	GAA Glu	CAC His	TAT Tyr	CAA Gln	CAG Gln	GTT Phe	TTT Phe	AAT Asn	GGT Gly	2784
		915					920					925				
TTG Leu	TCA Ser	ATA Ile	CTT Leu	GCC Ala	TCA Ser	TTT Phe	GCA Ala	TCC Ser	AGT Ser	GAT Asp	GAT Asp	CCA Pro	TAT Tyr	CAG Gln	GGC Gly	2832
	930					935					940					
CCG Pro	AGA Arg	GAT Asp	ATC Ile	ATA Ile	TCA Ser	CCT Pro	GAA Glu	CTG Leu	ATG Met	GCT Ala	GAT Asp	TAT Tyr	TTA Leu	CAA Gln	CCC Pro	2880
	945				950					955					960	
AAA Lys	TTG Leu	TTG Leu	GGC Gly	ATT Ile	TTG Leu	GCT Ala	TTT Phe	TTT Phe	AAC Asn	ATG Met	CAG Gln	TTA Leu	CTG Leu	AGC Ser	TCT Ser	2928
				965					970					975		
AGT Ser	GTT Val	GGC Gly	ATT Ile	GAA Glu	GAT Asp	AAG Lys	AAA Lys	ATG Met	GCC Ala	TTG Leu	AAC Asn	AGT Ser	TTG Leu	ATG Met	TCT Ser	2976
			980					985					990			
TTG Leu	ATG Met	AAG Lys	TTA Leu	ATG Met	GGA Gly	CCC Pro	AAA Lys	CAT His	GTC Val	AGT Ser	TCT Ser	GTG Val	AGG Arg	GTG Val	AAG Lys	3024
		995					1000						1005			
ATG Met	ATG Met	ACC Thr	ACA Thr	CTG Leu	AGA Arg	ACT Thr	GGC Gly	CTT Leu	CGA Arg	TTC Phe	AAG Lys	GAT Asp	GAT Asp	TTT Phe	CCT Pro	3072
	1010					1015					1020					
GAA Glu	TTG Leu	TGT Cys	TGC Cys	AGA Arg	GCT Ala	TGG Trp	GAC Asp	TGC Cys	TTT Phe	GTT Val	CGC Arg	TGC Cys	CTG Leu	GAT Asp	CAT His	3120
	1025				1030					1035					1040	



-83-

GCT TGT CTG GGC TCC CTT CTC AGT CAT GTA ATA GTA GCT TTG TTA CCT Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu Pro 1045 1050 1055	3168
CTT ATA CAC ATC CAG CCT AAA GAA ACT GCA GCT ATC TTC CAC TAC CTC Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr Leu 1060 1065 1070	3216
ATA ATT GAA AAC AGG GAT GCT GTG CAA GAT TTT CTT CAT GAA ATA TAT Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile Tyr 1075 1080 1085	3264
TTT TTA CCT GAT CAT CCA GAA TTA AAA AAG ATA AAA GCC GTT CTC CAG Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu Gln 1090 1095 1100	3312
GAA TAC AGA AAG GAG ACC TCT GAG AGC ACT GAT CTT CAG ACA ACT CTT Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr Leu 1105 1110 1115 1120	3360
CAG CTC TCT ATG AAG GCC ATT CAA CAT GAA AAT GTC GAT GTT CGT ATT Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg Ile 1125 1130 1135	3408
CAT GCT CTT ACA AGC TTG AAG GAA ACC TTG TAT AAA AAT CAG GAA AAA His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu Lys 1140 1145 1150	3456
CTG ATA AAG TAT GCA ACA GAC AGT GAA ACA GTA GAA CCT ATT ATC TCA Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile Ser 1155 1160 1165	3504
CAG TTG GTG ACA GTG CTT TTG AAA GGT TGC CAA GAT GCA AAC TCT CAA Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser Gln 1170 1175 1180	3552
GCT CGG TTG CTC TGT GGG GAA TGT TTA GGG GAA TTG GGG GCG ATA GAT Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile Asp 1185 1190 1195 1200	3600
CCA GGT CGA TTA GAT TTC TCA ACA ACT GAA ACT CAA GGA AAA GAT TTT Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp Phe 1205 1210 1215	3648
ACA TTT GTG ACT GGA GTA GAA GAT TCA AGC TTT GCC TAT GGA TTA TTG Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu Leu 1220 1225 1230	3696
ATG GAG CTA ACA AGA GCT TAC CTT GCG TAT GCT GAT AAT AGC CGA GCT Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg Ala 1235 1240 1245	3744
CCA GAT TCA GCT GCC TAT GCC ATT CAG GAG TTG CTT TCT ATT TAT GAC Pro Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr Asp 1250 1255 1260	3792
TGT AGA GAG ATG GAG ACC AAC GGC CCA GGT CAC CAA TTG TGG AGG AGA Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg Arg 1265 1270 1275 1280	3840
TTT CCT GAG CAT GTT CGG GAA ATA CTA GAA CCT CAT CTA AAT ACC AGA Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr Arg 1285 1290 1295	3888
TAC AAG AGT TCT CAG AAG TCA ACC GAT TGG TCT GGA GTA AAG AAG CCA Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys Pro 1300 1305 1310	3936

-84-

ATT TAC TTA AGT AAA TTG GGT AGT AAC TTT GCA GAA TGG TCA GCA TCT Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala Ser 1315 1320 1325	3984
TGG GCA GGT TAT CTT ATT ACA AAG GTT CGA CAT GAT CTT GCC AGT AAA Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser Lys 1330 1335 1340	4032
ATT TTC ACC TGC TGT AGC ATT ATG ATG AAG CAT GAT TTC AAA GTG ACC Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val Thr 1345 1350 1355 1360	4080
ATC TAT CTT CTT CCA CAT ATT CTG GTG TAT GTC TTA CTG GGT TGT AAT Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys Asn 1365 1370 1375	4128
CAA GAA GAT CAG CAG GAG GTT TAT GCA GAA ATT ATG GCA GTT CTA AAG Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu Lys 1380 1385 1390	4176
CAT GAC GAT CAG CAT ACC ATA AAT ACC CAA GAC ATT GCA TCT GAT CTG His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp Leu 1395 1400 1405	4224
TGT CAA CTC AGT ACA CAG ACT GTG TTC TCC ATG CTT GAC CAT CTC ACA Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu Thr 1410 1415 1420	4272
CAG TGG GCA AGG CAC AAA TTT CAG GCA CTG AAA GCT GAG AAA TGT CCA Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys Pro 1425 1430 1435 1440	4320
CAC AGC AAA TCA AAC AGA AAT AAG GTA GAC TCA ATG GTA TCT ACT GTG His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr Val 1445 1450 1455	4368
GAT TAT GAA GAC TAT CAG AGT GTA ACC CGT TTT CTA GAC CTC ATA CCC Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile Pro 1460 1465 1470	4416
CAG GAT ACT CTG GCA GTA GCT TCC TTT CGC TCC AAA GCA TAC ACA CGA Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr Arg 1475 1480 1485	4464
GCT GTA ATG CAC TTT GAA TCA TTT ATT ACA GAA AAG AAG CAA AAT ATT Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn Ile 1490 1495 1500	4512
CAG GAA CAT CTT GGA TTT TTA CAG AAA TTG TAT GCT GCT ATG CAT GAA Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His Glu 1505 1510 1515 1520	4560
CCT GAT GGA GTG TCC GGA GTC AGT GCA ATT AGA AAG GCA GAA CCA TCT Pro Asp Gly Val Ser Gly Val Ser Ala Ile Arg Lys Ala Glu Pro Ser 1525 1530 1535	4608
CTA AAA GAA CAG ATC CTT GAA CAT GAA AGC CTT GGC TTG CTG AGG GAT Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg Asp 1540 1545 1550	4656
GCC ACT GCT TGT TAT GAC AGG GCT ATT CAG CTA GAA CCA GAC CAG ATC Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln Ile 1555 1560 1565	4704
ATT CAT TAC CAT GGT GTA GTA AAG TCC ATG TTA GGT CTT GGT CAG CTG Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln Leu 1570 1575 1580	4752

-85-

TCT ACT GTT ATC ACT CAG GTG AAT GGA GTG CAT GCT AAC AGG TCC GAG Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser Glu 1585 1590 1595 1600	4800
TGG ACA GAT GAA TTA AAC ACG TAC AGA GTG GAA GCA GCT TGG AAA TTG Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys Leu 1605 1610 1615	4848
TCA CAG TGG GAT TTG GTG GAA AAC TAT TTG GCA GCA GAT GGA AAA TCT Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys Ser 1620 1625 1630	4896
ACA ACA TGG AGT GTC AGA CTG GGA CAG CTA TTA TTA TCA GCC AAA AAA Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys Lys 1635 1640 1645	4944
AGA GAT ATC ACA GCT TTT TAT GAC TCA CTG AAA CTA GTG AGA GCA GAA Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala Glu 1650 1655 1660	4992
CAA ATT GTA CCT CTT TCA GCT GCA AGC TTT GAA AGA GGC TCC TAC CAA Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr Gln 1665 1670 1675 1680	5040
CGA GGA TAT GAA TAT ATT GTG AGA TTG CAC ATG TTA TGT GAG TTG GAG Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu Glu 1685 1690 1695	5088
CAT AGC ATC AAA CCA CTT TTC CAG CAT TCT CCA GGT GAC AGT TCT CAA His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser Gln 1700 1705 1710	5136
GAA GAT TCT CTA AAC TGG GTA GCT CGA CTA GAA ATG ACC CAG AAT TCC Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn Ser 1715 1720 1725	5184
TAC AGA GCC AAG GAG CCT ATC CTG GCT CTC CGG AGG GCT TTA CTA AGC Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu Ser 1730 1735 1740	5232
CTC AAC AAA AGA CCA GAT TAC AAT GAA ATG GTT GGA GAA TGC TGG CTG Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp Leu 1745 1750 1755 1760	5280
CAG AGT GCC AGG GTA GCT AGA AAG GCT GGT CAC CAC CAG ACA GCC TAC Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala Tyr 1765 1770 1775	5328
AAT GCT CTC CTT AAT GCA GGG GAA TCA CGA CTC GCT GAA CTG TAC GTG Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr Val 1780 1785 1790	5376
GAA AGG GCA AAG TGG CTC TGG TCC AAG GGT GAT GTT CAC CAG GCA CTA Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala Leu 1795 1800 1805	5424
ATT GTT CTT CAA AAA GGT GTT GAA TTA TGT TTT CCT GAA AAT GAA ACC Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu Thr 1810 1815 1820	5472
CCA CCT GAG GGT AAG AAC ATG TTA ATC CAT GGT CGA GCT ATG CTA CTA Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu Leu 1825 1830 1835 1840	5520
GTG GGC CGA TTT ATG GAA GAA ACA GCT AAC TTT GAA AGC AAT GCA ATT Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala Ile 1845 1850 1855	5568

-86-

ATG AAA AAA TAT AAG GAT GTG ACC GCG TGC CTG CCA GAA TGG GAG GAT Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu Asp 1860 1865 1870	5616
GGG CAT TTT TAC CTT GCC AAG TAC TAT GAC AAA TTG ATG CCC ATG GTC Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met Val 1875 1880 1885	5664
ACA GAC AAC AAA ATG GAA AAG CAA GGT GAT CTC ATC CGG TAT ATA GTT Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile Val 1890 1895 1900	5712
CTT CAT TTT GGC AGA TCT CTA CAA TAT GGA AAT CAG TTC ATA TAT CAG Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr Gln 1905 1910 1915 1920	5760
TCA ATG CCA CGA ATG TTA ACT CTA TGG CTT GAT TAT GGT ACA AAG GCA Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys Ala 1925 1930 1935	5808
TAT GAA TGG GAA AAA GCT GGC CGC TCC GAT CGT GTA CAA ATG AGG AAT Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg Asn 1940 1945 1950	5856
GAT TTG GGT AAA ATA AAC AAG GTT ATC ACA GAG CAT ACA AAC TAT TTA Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr Leu 1955 1960 1965	5904
GCT CCA TAT CAA TTT TTG ACT GCT TTT TCA CAA TTG ATC TCT CGA ATT Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg Ile 1970 1975 1980	5952
TGT CAT TCT CAC GAT GAA GTT TTT GTT GTG CTT GAT GGA AAT AAT AGC Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn Ser 1985 1990 1995 2000	6000
CAA GTA TTT CTA GCC TAT CCT CAA CAA GCA ATG TGG ATG ATG ACA GCT Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr Ala 2005 2010 2015	6048
GTG TCA AAG TCA TCT TAT CCC ATG CGT GTG AAC AGA TGC AAG GAA ATC Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu Ile 2020 2025 2030	6096
CTC AAT AAA GCT ATT CAT ATG AAA AAA TCC TTA GAG AAG TTT GTT GGA Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val Gly 2035 2040 2045	6144
GAT GCA ACT CGC CTA ACA GAT AAG CTT CTA GAA TTG TGC AAT AAA CCG Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys Pro 2050 2055 2060	6192
GTT GAT GGA AGT AGT TCC ACA TTA AGC ATG AGC ACT CAT TTT AAA ATG Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys Met 2065 2070 2075 2080	6240
CTT AAA AAG CTG GTA GAA GAA GCA ACA TTT AGT GAA ATC CTC ATT CCT Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile Pro 2085 2090 2095	6288
CTA CAA TCA GTC ATG ATA CCT ACA CTT CCA TCA ATT CTG GGT ACC CAT Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr His 2100 2105 2110	6336
GCT AAC CAT GCT AGC CAT GAA CCA TTT CCT GGA CAT TGG GCC TAT ATT Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr Ile 2115 2120 2125	6384

-87-

GCA GGG TTT GAT GAT ATG GTG GAA ATT CTT GCT TCT CTT CAG AAA CCA Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys Pro 2130 2135 2140	6432
AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG TTC TAC ATC ATG ATG Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met Met 2145 2150 2155 2160	6480
TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT AGA CTA ATG GAA TTC Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu Phe 2165 2170 2175	6528
AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT GCA GAG TCT CGT AGA Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg Arg 2180 2185 2190	6576
AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT CCA CTA AAT GAT GAA Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp Glu 2195 2200 2205	6624
TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT GGT TTG AGA CCT ATT Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro Ile 2210 2215 2220	6672
CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT ATG ACA GGA AAA GAA Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys Glu 2225 2230 2235 2240	6720
CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT TTA TCT GAA AAA CTC Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys Leu 2245 2250 2255	6768
AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT CCT CCT ATT TTT CAT Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro Ile Phe His 2260 2265 2270	6816
GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA TCA TGG TAC AGT AGT Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser Ser 2275 2280 2285	6864
AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA ATG GTT GGT TAT Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly Tyr 2290 2295 2300	6912
ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT CTC TTT GAT TCT Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp Ser 2305 2310 2315 2320	6960
TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT CTT TTC AAT AAG Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn Lys 2325 2330 2335	7008
GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT CGC CTG ACT CAT Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr His 2340 2345 2350	7056
AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG GGT CTT TTT CGA Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe Arg 2355 2360 2365	7104
AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT CAG CGA GAG CCT Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu Pro 2370 2375 2380	7152
TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT CTT GTG GAA TGG Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu Trp 2385 2390 2395 2400	7200

-88-

AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG AAT GAA ACT GGA	7248
Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr Gly	
2405 2410 2415	
GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT GAC ATT GAG CAG	7296
Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu Gln	
2420 2425 2430	
CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG ACA GGA CTG CCG	7344
Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu Pro	
2435 2440 2445	
TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA GAA GCT ACT GAT	7392
Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr Asp	
2450 2455 2460	
GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT CCA TAT ATG	7437
Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met	
2465 2470 2475	
TGA	7440

## (2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2479 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser	
1 5 10 15	
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu	
20 25 30	
Met Ser Met Gln Lys Leu Glu Phe Ile Glu Val Thr Leu Leu Thr Val	
35 40 45	
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu	
50 55 60	
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile	
65 70 75 80	
Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly	
85 90 95	
Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu	
100 105 110	
Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu	
115 120 125	
Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu	
130 135 140	
Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys	
145 150 155 160	
Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp	
165 170 175	

-89-

Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe  
 180 185 190  
 Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val  
 195 200 205  
 Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly Ile Glu Val Asp  
 210 215 220  
 Ala Glu Tyr Leu Leu Gly Pro Leu Tyr Ala Ala Leu Lys Met Glu Ser  
 225 230 235 240  
 Met Glu Ile Ile Glu Glu Ile Gln Cys Gln Thr Gln Gln Glu Asn Leu  
 245 250 255  
 Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg Arg Leu Ser Ser  
 260 265 270  
 Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr Glu Glu Ile Lys  
 275 280 285  
 His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser Ala Leu Lys Gln  
 290 295 300  
 Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser Gly Leu Lys Asn  
 305 310 315 320  
 Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val Leu Gln Leu Thr  
 325 330 335  
 Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met Asn Cys Arg Thr  
 340 345 350  
 Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro Ser Val Val Ile  
 355 360 365  
 Thr Trp Met Ser Leu Asp Phe Tyr Thr Thr Val Leu Lys Ser Cys Arg  
 370 375 380  
 Arg Leu Leu Glu Ser Val Gln Lys Arg Thr Gly Gly Asn Ile Asp Lys  
 385 390 395 400  
 Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met Gln Val Asn Ser Ser  
 405 410 415  
 Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly Met Leu Ser Leu Pro  
 420 425 430  
 Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu Lys Leu Thr Thr Phe  
 435 440 445  
 Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile Ser Asp Ser Tyr Ser  
 450 455 460  
 Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu Thr Leu Phe Pro Arg  
 465 470 475 480  
 Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr Asn Trp Ala Leu Gln  
 485 490 495  
 Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe Ile  
 500 505 510  
 Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu Ile  
 515 520 525

-90-

Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala Ser  
 530 535 540  
 Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu Thr  
 545 550 555 560  
 Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu Phe  
 565 570 575  
 Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser Gln  
 580 585 590  
 Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe Leu Leu Lys Lys Lys  
 595 600 605  
 Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp Asn Leu His His Leu  
 610 615 620  
 Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr Asp Val Lys Ala Val  
 625 630 635 640  
 Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro Asp Lys Asp Val Arg  
 645 650 655  
 Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu Glu Ser Leu Asp Ser  
 660 665 670  
 Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu Arg Met Lys Glu Ala  
 675 680 685  
 Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu Leu Lys Asp Thr Leu  
 690 695 700  
 Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala Lys Gly Asp Leu Val  
 705 710 715 720  
 Pro Phe Ala Leu Leu His Leu Leu His Cys Leu Leu Ser Lys Ser Ala  
 725 730 735  
 Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg Ala Leu Val Ala Ala  
 740 745 750  
 Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln Tyr Lys Lys Pro Ile  
 755 760 765  
 Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser Gln Met Thr Ala Leu  
 770 775 780  
 Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val Ala  
 785 790 795 800  
 His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn Val  
 805 810 815  
 Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln Val  
 820 825 830  
 Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala Leu  
 835 840 845  
 Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile Leu  
 850 855 860  
 Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys Ser  
 865 870 875 880



Lys	Asp	Glu	Leu	Glu	Arg	Ala	Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr	Glu	885	890	895
Ile	Glu	Leu	Gly	Ser	Leu	Leu	Arg	Gln	Asp	Phe	Gln	Gly	Leu	His	Asn	900	905	910
Glu	Leu	Leu	Leu	Arg	Ile	Gly	Glu	His	Tyr	Gln	Gln	Val	Phe	Asn	Gly	915	920	925
Leu	Ser	Ile	Leu	Ala	Ser	Phe	Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln	Gly	930	935	940
Pro	Arg	Asp	Ile	Ile	Ser	Pro	Glu	Leu	Met	Ala	Asp	Tyr	Leu	Gln	Pro	945	950	955
Lys	Leu	Leu	Gly	Ile	Leu	Ala	Phe	Phe	Asn	Met	Gln	Leu	Leu	Ser	Ser	965	970	975
Ser	Val	Gly	Ile	Glu	Asp	Lys	Lys	Met	Ala	Leu	Asn	Ser	Leu	Met	Ser	980	985	990
Leu	Met	Lys	Leu	Met	Gly	Pro	Lys	His	Val	Ser	Ser	Val	Arg	Val	Lys	995	1000	1005
Met	Met	Thr	Thr	Leu	Arg	Thr	Gly	Leu	Arg	Phe	Lys	Asp	Asp	Phe	Pro	1010	1015	1020
Glu	Leu	Cys	Cys	Arg	Ala	Trp	Asp	Cys	Phe	Val	Arg	Cys	Leu	Asp	His	1025	1030	1035
Ala	Cys	Leu	Gly	Ser	Leu	Leu	Ser	His	Val	Ile	Val	Ala	Leu	Leu	Pro	1045	1050	1055
Leu	Ile	His	Ile	Gln	Pro	Lys	Glu	Thr	Ala	Ala	Ile	Phe	His	Tyr	Leu	1060	1065	1070
Ile	Ile	Glu	Asn	Arg	Asp	Ala	Val	Gln	Asp	Phe	Leu	His	Glu	Ile	Tyr	1075	1080	1085
Phe	Leu	Pro	Asp	His	Pro	Glu	Leu	Lys	Lys	Ile	Lys	Ala	Val	Leu	Gln	1090	1095	1100
Glu	Tyr	Arg	Lys	Glu	Thr	Ser	Glu	Ser	Thr	Asp	Leu	Gln	Thr	Thr	Leu	1105	1110	1115
Gln	Leu	Ser	Met	Lys	Ala	Ile	Gln	His	Glu	Asn	Val	Asp	Val	Arg	Ile	1125	1130	1135
His	Ala	Leu	Thr	Ser	Leu	Lys	Glu	Thr	Leu	Tyr	Lys	Asn	Gln	Glu	Lys	1140	1145	1150
Leu	Ile	Lys	Tyr	Ala	Thr	Asp	Ser	Glu	Thr	Val	Glu	Pro	Ile	Ile	Ser	1155	1160	1165
Gln	Leu	Val	Thr	Val	Leu	Leu	Lys	Gly	Cys	Gln	Asp	Ala	Asn	Ser	Gln	1170	1175	1180
Ala	Arg	Leu	Leu	Cys	Gly	Glu	Cys	Leu	Gly	Glu	Leu	Gly	Ala	Ile	Asp	1185	1190	1195
Pro	Gly	Arg	Leu	Asp	Phe	Ser	Thr	Thr	Glu	Thr	Gln	Gly	Lys	Asp	Phe	1205	1210	1215
Thr	Phe	Val	Thr	Gly	Val	Glu	Asp	Ser	Ser	Phe	Ala	Tyr	Gly	Leu	Leu	1220	1225	1230

-92-

Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg Ala  
1235 1240 1245

Pro Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr Asp  
1250 1255 1260

Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg Arg  
1265 1270 1275 1280

Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr Arg  
1285 1290 1295

Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys Pro  
1300 1305 1310

Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala Ser  
1315 1320 1325

Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser Lys  
1330 1335 1340

Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val Thr  
1345 1350 1355 1360

Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys Asn  
1365 1370 1375

Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu Lys  
1380 1385 1390

His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp Leu  
1395 1400 1405

Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu Thr  
1410 1415 1420

Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys Pro  
1425 1430 1435 1440

His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr Val  
1445 1450 1455

Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile Pro  
1460 1465 1470

Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr Arg  
1475 1480 1485

Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn Ile  
1490 1495 1500

Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His Glu  
1505 1510 1515 1520

Pro Asp Gly Val Ser Gly Val Ser Ala Ile Arg Lys Ala Glu Pro Ser  
1525 1530 1535

Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg Asp  
1540 1545 1550

Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln Ile  
1555 1560 1565

Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln Leu  
1570 1575 1580

-93-

Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser Glu  
 1585 1590 1595 1600  
 Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys Leu  
 1605 1610 1615  
 Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys Ser  
 1620 1625 1630  
 Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys Lys  
 1635 1640 1645  
 Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala Glu  
 1650 1655 1660  
 Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr Gln  
 1665 1670 1675 1680  
 Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu Glu  
 1685 1690 1695  
 His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser Gln  
 1700 1705 1710  
 Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn Ser  
 1715 1720 1725  
 Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu Ser  
 1730 1735 1740  
 Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp Leu  
 1745 1750 1755 1760  
 Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala Tyr  
 1765 1770 1775  
 Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr Val  
 1780 1785 1790  
 Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala Leu  
 1795 1800 1805  
 Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu Thr  
 1810 1815 1820  
 Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu Leu  
 1825 1830 1835 1840  
 Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala Ile  
 1845 1850 1855  
 Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu Asp  
 1860 1865 1870  
 Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met Val  
 1875 1880 1885  
 Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile Val  
 1890 1895 1900  
 Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr Gln  
 1905 1910 1915 1920  
 Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys Ala  
 1925 1930 1935

-94-

Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg Asn  
 1940 1945 1950  
 Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr Leu  
 1955 1960 1965  
 Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg Ile  
 1970 1975 1980  
 Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn Ser  
 1985 1990 1995 2000  
 Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr Ala  
 2005 2010 2015  
 Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu Ile  
 2020 2025 2030  
 Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val Gly  
 2035 2040 2045  
 Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys Pro  
 2050 2055 2060  
 Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys Met  
 2065 2070 2075 2080  
 Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile Pro  
 2085 2090 2095  
 Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr His  
 2100 2105 2110  
 Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr Ile  
 2115 2120 2125  
 Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys Pro  
 2130 2135 2140  
 Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met Met  
 2145 2150 2155 2160  
 Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu Phe  
 2165 2170 2175  
 Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg Arg  
 2180 2185 2190  
 Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp Glu  
 2195 2200 2205  
 Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro Ile  
 2210 2215 2220  
 Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys Glu  
 2225 2230 2235 2240  
 Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys Leu  
 2245 2250 2255  
 Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro Ile Phe His  
 2260 2265 2270  
 Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser Ser  
 2275 2280 2285

-95-

Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly Tyr  
 2290 2295 2300  
 Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp Ser  
 2305 2310 2315 2320  
 Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn Lys  
 2325 2330 2335  
 Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr His  
 2340 2345 2350  
 Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe Arg  
 2355 2360 2365  
 Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu Pro  
 2370 2375 2380  
 Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu Trp  
 2385 2390 2395 2400  
 Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr Gly  
 2405 2410 2415  
 Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu Gln  
 2420 2425 2430  
 Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu Pro  
 2435 2440 2445  
 Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr Asp  
 2450 2455 2460  
 Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2465 2470 2475

## (2) INFORMATION FOR SEQ ID NO:19:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 23 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: Primer oDH26

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

TGGTTTCTGA GAACATTCCC TGA

23

## (2) INFORMATION FOR SEQ ID NO:20:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 9 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: peptide

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: FLAG tag

-96-

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

Met Asp Tyr Lys Asp Asp Asp Asp Lys  
1 5

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 20 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:  
(B) CLONE: Primer 279-3

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

TGGATGATGA CAGCTGTGTC

20

(2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 20 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

- (vii) IMMEDIATE SOURCE:  
(B) CLONE: Primer 279-6

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

TGTAGTCGCT GCTCAATGTC

20

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 7624 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:  
(A) NAME/KEY: CDS  
(B) LOCATION: 333..7562

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

CTTGTGAAGA GAATGTTTTA CACTCTTGTT AGTGAAGTTT ATTCTTTAAA AGTCAATCGT

60

CAAGGATTTA GCAAATGAAT TAGCACTTCG GATATACTTG TTTATTTAAT ATCTTTTTTG

120

-97-

TTTATTTCAA	AGAATTCAGT	AATTGGATCA	TAACGAGACT	TCTGCGGATT	GCAGCAACTC	180										
CCTCCTGTCA	TTTGTTACAC	AAGAAAATCT	GTGAAGTCAT	CTGTTTCATTA	TTATTTCTTT	240										
TTAAAAGCAA	GAGTCCTGCT	ATTTTTGGGG	TACTCACAAA	AGAATTATTA	CAACTTTTGTG	300										
AAGACTTGGT	TTACCTCCAT	AGAAGAAATG	TG	ATG	GGT	CAT	GCT	GTG	GAA	TGG	353					
				Met	Gly	His	Ala	Val	Glu	Trp						
				1				5								
CCA	GTG	GTC	ATG	AGC	CGA	TTT	TTA	AGT	CAA	TTA	GAT	GAA	CAC	ATG	GGA	401
Pro	Val	Val	Met	Ser	Arg	Phe	Leu	Ser	Gln	Leu	Asp	Glu	His	Met	Gly	
	10						15				20					
TAT	TTA	CAA	TCA	GCT	CCT	TTG	CAG	TTG	ATG	AGT	ATG	CAA	AAT	TTA	GAA	449
Tyr	Leu	Gln	Ser	Ala	Pro	Leu	Gln	Leu	Met	Ser	Met	Gln	Asn	Leu	Glu	
	25					30					35					
TTT	ATT	GAA	GTC	ACT	TTA	TTA	ATG	GTT	CTT	ACT	CGT	ATT	ATT	GCA	ATT	497
Phe	Ile	Glu	Val	Thr	Leu	Leu	Met	Val	Leu	Thr	Arg	Ile	Ile	Ala	Ile	
	40				45					50					55	
GTG	TTT	TTT	AGA	AGG	CAA	GAA	CTC	TTA	CTT	TGG	CAG	ATA	GGT	TGT	GTT	545
Val	Phe	Phe	Arg	Arg	Gln	Glu	Leu	Leu	Leu	Trp	Gln	Ile	Gly	Cys	Val	
				60					65					70		
CTG	CTA	GAG	TAT	GGT	AGT	CCA	AAA	ATT	AAA	TCC	CTA	GCA	ATT	AGC	TTT	593
Leu	Leu	Glu	Tyr	Gly	Ser	Pro	Lys	Ile	Lys	Ser	Leu	Ala	Ile	Ser	Phe	
			75					80					85			
TTA	ACA	GAA	CTT	TTT	CAG	CTT	GGA	GGA	CTA	CCA	GCA	CAA	CCA	GCT	AGC	641
Leu	Thr	Glu	Leu	Phe	Gln	Leu	Gly	Gly	Leu	Pro	Ala	Gln	Pro	Ala	Ser	
	90						95					100				
ACT	TTT	TTC	AGC	TCA	TTT	TTG	GAA	TTA	TTA	AAA	CAC	CTT	GTA	GAA	ATG	689
Thr	Phe	Phe	Ser	Ser	Phe	Leu	Glu	Leu	Leu	Lys	His	Leu	Val	Glu	Met	
	105					110					115					
GAT	ACT	GAC	CAA	TTG	AAA	CTC	TAT	GAA	GAG	CCA	TTA	TCA	AAG	CTG	ATA	737
Asp	Thr	Asp	Gln	Leu	Lys	Leu	Tyr	Glu	Glu	Pro	Leu	Ser	Lys	Leu	Ile	
	120				125					130					135	
AAG	ACA	CTA	TTT	CCC	TTT	GAA	GCA	GAA	GCT	TAT	AGA	AAT	ATT	GAA	CCT	785
Lys	Thr	Leu	Phe	Pro	Phe	Glu	Ala	Glu	Ala	Tyr	Arg	Asn	Ile	Glu	Pro	
				140					145					150		
GTC	TAT	TTA	AAT	ATG	CTG	CTG	GAA	AAA	CTC	TGT	GTC	ATG	TTT	GAA	GAC	833
Val	Tyr	Leu	Asn	Met	Leu	Leu	Glu	Lys	Leu	Cys	Val	Met	Phe	Glu	Asp	
			155					160					165			
GGT	GTG	CTC	ATG	CGG	CTT	AAG	TCT	GAT	TTG	CTA	AAA	GCA	GCT	TTG	TGC	881
Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	Leu	Leu	Lys	Ala	Ala	Leu	Cys	
		170					175					180				
CAT	TTA	CTG	CAG	TAT	TTC	CTT	AAA	TTT	GTG	CCA	GCT	GGG	TAT	GAA	TCT	929
His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	Val	Pro	Ala	Gly	Tyr	Glu	Ser	
	185					190					195					
GCT	TTA	CAA	GTC	AGG	AAG	GTC	TAT	GTG	AGA	AAT	ATT	TGT	AAA	GCT	CTT	977
Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	Arg	Asn	Ile	Cys	Lys	Ala	Leu	
	200				205					210					215	
TTG	GAT	GTG	CTT	GGA	ATT	GAG	GTA	GAT	GCA	GAG	TAC	TTG	TTG	GGC	CCA	1025
Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	Ala	Glu	Tyr	Leu	Leu	Gly	Pro	
				220					225					230		

-98-

CTT TAT GCA GCT TTG AAA ATG GAA AGT ATG GAA ATC ATT GAG GAG ATT	1073
Leu Tyr Ala Ala Leu Lys Met Glu Ser Met Glu Ile Ile Glu Glu Ile	
235 240 245	
CAA TGC CAA ACT CAA CAG GAA AAC CTC AGC AGT AAT AGT GAT GGA ATA	1121
Gln Cys Gln Thr Gln Gln Glu Asn Leu Ser Ser Asn Ser Asp Gly Ile	
250 255 260	
TCA CCC AAA AGG CGT CGT CTC AGC TCG TCT CTA AAC CCT TCT AAA AGA	1169
Ser Pro Lys Arg Arg Arg Leu Ser Ser Ser Leu Asn Pro Ser Lys Arg	
265 270 275	
GCA CCA AAA CAG ACT GAG GAA ATT AAA CAT GTG GAC ATG AAC CAA AAG	1217
Ala Pro Lys Gln Thr Glu Glu Ile Lys His Val Asp Met Asn Gln Lys	
280 285 290 295	
AGC ATA TTA TGG AGT GCA CTG AAA CAG AAA GCT GAA TCC CTT CAG ATT	1265
Ser Ile Leu Trp Ser Ala Leu Lys Gln Lys Ala Glu Ser Leu Gln Ile	
300 305 310	
TCC CTT GAA TAC AGT GGC CTA AAG AAT CCT GTT ATT GAG ATG TTA GAA	1313
Ser Leu Glu Tyr Ser Gly Leu Lys Asn Pro Val Ile Glu Met Leu Glu	
315 320 325	
GGA ATT GCT GTT GTC TTA CAA CTG ACT GCT CTG TGT ACT GTT CAT TGT	1361
Gly Ile Ala Val Val Leu Gln Leu Thr Ala Leu Cys Thr Val His Cys	
330 335 340	
TCT CAT CAA AAC ATG AAC TGC CGT ACT TTC AAG GAC TGT CAA CAT AAA	1409
Ser His Gln Asn Met Asn Cys Arg Thr Phe Lys Asp Cys Gln His Lys	
345 350 355	
TCC AAG AAG AAA CCT TCT GTA GTG ATA ACT TGG ATG TCA TTG GAT TTT	1457
Ser Lys Lys Lys Pro Ser Val Val Ile Thr Trp Met Ser Leu Asp Phe	
360 365 370 375	
TAC ACA AAA GTG CTT AAG AGC TGT AGA AGT TTG TTA GAA TCT GTT CAG	1505
Tyr Thr Lys Val Leu Lys Ser Cys Arg Ser Leu Leu Glu Ser Val Gln	
380 385 390	
AAA CTG GAC CTG GAG GCA ACC ATT GAT AAG GTG GTG AAA ATT TAT GAT	1553
Lys Leu Asp Leu Glu Ala Thr Ile Asp Lys Val Val Lys Ile Tyr Asp	
395 400 405	
GCT TTG ATT TAT ATG CAA GTA AAC AGT TCA TTT GAA GAT CAT ATC CTG	1601
Ala Leu Ile Tyr Met Gln Val Asn Ser Ser Phe Glu Asp His Ile Leu	
410 415 420	
GAA GAT TTA TGT GGA ATG CTC TCA CTT CCA TGG ATT TAT TCC CAT TCT	1649
Glu Asp Leu Cys Gly Met Leu Ser Leu Pro Trp Ile Tyr Ser His Ser	
425 430 435	
GAT GAT GGC TGT TTA AAG TTG ACC ACA TTT GCC GCT AAT CTT CTA ACA	1697
Asp Asp Gly Cys Leu Lys Leu Thr Thr Phe Ala Ala Asn Leu Leu Thr	
440 445 450 455	
TTA AGC TGT AGG ATT TCA GAT AGC TAT TCA CCA CAG GCA CAA TCA CGA	1745
Leu Ser Cys Arg Ile Ser Asp Ser Tyr Ser Pro Gln Ala Gln Ser Arg	
460 465 470	
TGT GTG TTT CTT CTG ACT CTG TTT CCA AGA AGA ATA TTC CTT GAG TGG	1793
Cys Val Phe Leu Leu Thr Leu Phe Pro Arg Arg Ile Phe Leu Glu Trp	
475 480 485	
AGA ACA GCA GTT TAC AAC TGG GCC CTG CAG AGC TCC CAT GAA GTA ATC	1841
Arg Thr Ala Val Tyr Asn Trp Ala Leu Gln Ser Ser His Glu Val Ile	
490 495 500	



-99-

CGG Arg 505	GCT Ala	AGT Ser	TGT Cys	GTT Val	AGT Ser	GGA Gly 510	TTT Phe	TTT Phe	ATC Ile	TTA Leu	TTG Leu 515	CAG Gln	CAG Gln	CAG Gln	AAT Asn	1889
TCT Ser 520	TGT Cys	AAC Asn	AGA Arg	GTT Val	CCC Pro 525	AAG Lys	ATT Ile	CTT Leu	ATA Ile	GAT Asp 530	AAA Lys	GTC Val	AAA Lys	GAT Asp	GAT Asp 535	1937
TCT Ser	GAC Asp	ATT Ile	GTC Val	AAG Lys 540	AAA Lys	GAA Glu	TTT Phe	GCT Ala	TCT Ser 545	ATA Ile	CTT Leu	GGT Gly	CAA Gln	CTT Leu 550	GTC Val	1985
TGT Cys	ACT Thr	CTT Leu	CAC His 555	GGC Gly	ATG Met	TTT Phe	TAT Tyr	CTG Leu 560	ACA Thr	AGT Ser	TCT Ser	TTA Leu	ACA Thr 565	GAA Glu	CCT Pro	2033
TTC Phe	TCT Ser	GAA Glu 570	CAC His	GGA Gly	CAT His	GTG Val	GAC Asp 575	CTC Leu	TTC Phe	TGT Cys	AGG Arg	AAC Asn 580	TTG Leu	AAA Lys	GCC Ala	2081
ACT Thr	TCT Ser 585	CAA Gln	CAT His	GAA Glu	TGT Cys	TCA Ser 590	TCT Ser	TCT Ser	CAA Gln	CTA Leu	AAA Lys 595	GCT Ala	TCT Ser	GTC Val	TGC Cys	2129
AAG Lys 600	CCA Pro	TTC Phe	CTT Leu	TTC Phe	CTA Leu 605	CTG Leu	AAA Lys	AAA Lys	AAA Lys	ATA Ile 610	CCT Pro	AGT Ser	CCA Pro	GTA Val	AAA Lys 615	2177
CTT Leu	GCT Ala	TTC Phe	ATA Ile	GAT Asp 620	AAT Asn	CTA Leu	CAT His	CAT His	CTT Leu 625	TGT Cys	AAG Lys	CAT His	CTT Leu	GAT Asp 630	TTT Phe	2225
AGA Arg	GAA Glu	GAT Asp	GAA Glu 635	ACA Thr	GAT Asp	GTA Val	AAA Lys	GCA Ala 640	GTT Val	CTT Leu	GGA Gly	ACT Thr	TTA Leu 645	TTA Leu	AAT Asn	2273
TTA Leu	ATG Met	GAA Glu 650	GAT Asp	CCA Pro	GAC Asp	AAA Lys	GAT Asp 655	GTT Val	AGA Arg	GTG Val	GCT Ala	TTT Phe 660	AGT Ser	GGA Gly	AAT Asn	2321
ATC Ile 665	AAG Lys	CAC His	ATA Ile	TTG Leu	GAA Glu	TCC Ser 670	TTG Leu	GAC Asp	TCT Ser	GAA Glu	GAT Asp 675	GGA Gly	TTT Phe	ATA Ile	AAG Lys	2369
GAG Glu 680	CTT Leu	TTT Phe	GTC Val	TTA Leu	AGA Arg 685	ATG Met	AAG Lys	GAA Glu	GCA Ala 690	TAT Tyr	ACA Thr	CAT His	GCC Ala	CAA Gln	ATA Ile 695	2417
TCA Ser	AGA Arg	AAT Asn	AAT Asn	GAG Glu 700	CTG Leu	AAG Lys	GAT Asp	ACC Thr	TTG Leu 705	ATT Ile	CTT Leu	ACA Thr	ACA Thr	GGG Gly 710	GAT Asp	2465
ATT Ile	GGA Gly	AGG Arg	GCC Ala 715	GCA Ala	AAA Lys	GGA Gly	GAT Asp 720	TTG Leu	GTA Val	CCA Pro	TTT Phe	GCA Ala 725	CTC Ala	TTA Leu	CAC His	2513
TTA Leu	TTG Leu	CAT His 730	TGT Cys	TTG Leu	TTA Leu	TCC Ser	AAG Lys 735	TCA Ser	GCA Ala	TCT Ser	GTC Val	TCT Ser 740	GGA Gly	GCA Ala	GCA Ala	2561
TAC Tyr	ACA Thr 745	GAA Glu	ATT Ile	AGA Arg	GCT Ala	CTG Leu 750	GTT Val	GCA Ala	GCT Ala	AAA Lys	AGT Ser 755	GTT Val	AAA Lys	CTG Leu	CAA Gln	2609
AGT Ser 760	TTT Phe	TTC Phe	AGC Ser	CAG Gln	TAT Tyr 765	AAG Lys	AAA Lys	CCC Pro	ATC Ile	TGT Cys 770	CAG Gln	TTT Phe	TTG Leu	GTA Val	GAA Glu 775	2657

-100-

TCC	CTT	CAC	TCT	AGT	CAG	ATG	ACA	GCA	CTT	CCG	AAT	ACT	CCA	TGC	CAG	2705
Ser	Leu	His	Ser	Ser	Gln	Met	Thr	Ala	Leu	Pro	Asn	Thr	Pro	Cys	Gln	
				780					785					790		
AAT	GCT	GAC	GTG	CGA	AAA	CAA	GAT	GTG	GCT	CAC	CAG	AGA	GAA	ATG	GCT	2753
Asn	Ala	Asp	Val	Arg	Lys	Gln	Asp	Val	Ala	His	Gln	Arg	Glu	Met	Ala	
			795					800					805			
TTA	AAT	ACG	TTG	TCT	GAA	ATT	GCC	AAC	GTT	TTC	GAC	TTT	CCT	GAT	CTT	2801
Leu	Asn	Thr	Leu	Ser	Glu	Ile	Ala	Asn	Val	Phe	Asp	Phe	Pro	Asp	Leu	
		810					815					820				
AAT	CGT	TTT	CTT	ACT	AGG	ACA	TTA	CAA	GTT	CTA	CTA	CCT	GAT	CTT	GCT	2849
Asn	Arg	Phe	Leu	Thr	Arg	Thr	Leu	Gln	Val	Leu	Leu	Pro	Asp	Leu	Ala	
	825					830					835					
GCC	AAA	GCA	AGC	CCT	GCA	GCT	TCT	GCT	CTC	ATT	CGA	ACT	TTA	GGA	AAA	2897
Ala	Lys	Ala	Ser	Pro	Ala	Ala	Ser	Ala	Leu	Ile	Arg	Thr	Leu	Gly	Lys	
840					845					850					855	
CAA	TTA	AAT	GTC	AAT	CGT	AGA	GAG	ATT	TTA	ATA	AAC	AAC	TTC	AAA	TAT	2945
Gln	Leu	Asn	Val	Asn	Arg	Arg	Glu	Ile	Leu	Ile	Asn	Asn	Phe	Lys	Tyr	
				860					865					870		
ATT	TTT	TCT	CAT	TTG	GTC	TGT	TCT	TGT	TCC	AAA	GAT	GAA	TTA	GAA	CGT	2993
Ile	Phe	Ser	His	Leu	Val	Cys	Ser	Cys	Ser	Lys	Asp	Glu	Leu	Glu	Arg	
			875					880					885			
GCC	CTT	CAT	TAT	CTG	AAG	AAT	GAA	ACA	GAA	ATT	GAA	CTG	GGG	AGC	CTG	3041
Ala	Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr	Glu	Ile	Glu	Leu	Gly	Ser	Leu	
		890					895					900				
TTG	AGA	CAA	GAT	TTC	CAA	GGA	TTG	CAT	AAT	GAA	TTA	TTG	CTG	CGT	ATT	3089
Leu	Arg	Gln	Asp	Phe	Gln	Gly	Leu	His	Asn	Glu	Leu	Leu	Leu	Arg	Ile	
	905					910					915					
GGA	GAA	CAC	TAT	CAA	CAG	GTT	TTT	AAT	GGT	TTG	TCA	ATA	CTT	GCC	TCA	3137
Gly	Glu	His	Tyr	Gln	Gln	Val	Phe	Asn	Gly	Leu	Ser	Ile	Leu	Ala	Ser	
920					925					930					935	
TTT	GCA	TCC	AGT	GAT	GAT	CCA	TAT	CAG	GGC	CCG	AGA	GAT	ATC	ATA	TCA	3185
Phe	Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln	Gly	Pro	Arg	Asp	Ile	Ile	Ser	
				940					945					950		
CCT	GAA	CTG	ATG	GCT	GAT	TAT	TTA	CAA	CCC	AAA	TTG	TTG	GGC	ATT	TTG	3233
Pro	Glu	Leu	Met	Ala	Asp	Tyr	Leu	Gln	Pro	Lys	Leu	Leu	Gly	Ile	Leu	
			955					960					965			
GCT	TTT	TTT	AAC	ATG	CAG	TTA	CTG	AGC	TCT	AGT	GTT	GGC	ATT	GAA	GAT	3281
Ala	Phe	Phe	Asn	Met	Gln	Leu	Leu	Ser	Ser	Ser	Val	Gly	Ile	Glu	Asp	
		970					975					980				
AAG	AAA	ATG	GCC	TTG	AAC	AGT	TTG	ATG	TCT	TTG	ATG	AAG	TTA	ATG	GGA	3329
Lys	Lys	Met	Ala	Leu	Asn	Ser	Leu	Met	Ser	Leu	Met	Lys	Leu	Met	Gly	
	985					990					995					
CCC	AAA	CAT	GTC	AGT	TCT	GTG	AGG	GTG	AAG	ATG	ATG	ACC	ACA	CTG	AGA	3377
Pro	Lys	His	Val	Ser	Ser	Val	Arg	Val	Lys	Met	Met	Thr	Thr	Leu	Arg	
1000						1005				1010					1015	
ACT	GGC	CTT	CGA	TTC	AAG	GAT	GAT	TTT	CCT	GAA	TTG	TGT	TGC	AGA	GCT	3425
Thr	Gly	Leu	Arg	Phe	Lys	Asp	Asp	Phe	Pro	Glu	Leu	Cys	Cys	Arg	Ala	
				1020					1025					1030		
TGG	GAC	TGC	TTT	GTT	CGC	TGC	CTG	GAT	CAT	GCT	TGT	CTG	GGC	TCC	CTT	3473
Trp	Asp	Cys	Phe	Val	Arg	Cys	Leu	Asp	His	Ala	Cys	Leu	Gly	Ser	Leu	
			1035					1040					1045			

-101-

CTC	AGT	CAT	GTA	ATA	GTA	GCT	TTG	TTA	CCT	CTT	ATA	CAC	ATC	CAG	CCT	3521
Leu	Ser	His	Val	Ile	Val	Ala	Leu	Leu	Pro	Leu	Ile	His	Ile	Gln	Pro	
		1050					1055					1060				
AAA	GAA	ACT	GCA	GCT	ATC	TTC	CAC	TAC	CTC	ATA	ATT	GAA	AAC	AGG	GAT	3569
Lys	Glu	Thr	Ala	Ala	Ile	Phe	His	Tyr	Leu	Ile	Ile	Glu	Asn	Arg	Asp	
	1065					1070					1075					
GCT	GTG	CAA	GAT	TTT	CTT	CAT	GAA	ATA	TAT	TTT	TTA	CCT	GAT	CAT	CCA	3617
Ala	Val	Gln	Asp	Phe	Leu	His	Glu	Ile	Tyr	Phe	Leu	Pro	Asp	His	Pro	
1080					1085					1090					1095	
GAA	TTA	AAA	AAG	ATA	AAA	GCC	GTT	CTC	CAG	GAA	TAC	AGA	AAG	GAG	ACC	3665
Glu	Leu	Lys	Lys	Ile	Lys	Ala	Val	Leu	Gln	Glu	Tyr	Arg	Lys	Glu	Thr	
				1100					1105					1110		
TCT	GAG	AGC	ACT	GAT	CTT	CAG	ACA	ACT	CTT	CAG	CTC	TCT	ATG	AAG	GCC	3713
Ser	Glu	Ser	Thr	Asp	Leu	Gln	Thr	Thr	Leu	Gln	Leu	Ser	Met	Lys	Ala	
			1115					1120					1125			
ATT	CAA	CAT	GAA	AAT	GTC	GAT	GTT	CGT	ATT	CAT	GCT	CTT	ACA	AGC	TTG	3761
Ile	Gln	His	Glu	Asn	Val	Asp	Val	Arg	Ile	His	Ala	Leu	Thr	Ser	Leu	
		1130					1135					1140				
AAG	GAA	ACC	TTG	TAT	AAA	AAT	CAG	GAA	AAA	CTG	ATA	AAG	TAT	GCA	ACA	3809
Lys	Glu	Thr	Leu	Tyr	Lys	Asn	Gln	Glu	Lys	Leu	Ile	Lys	Tyr	Ala	Thr	
	1145					1150					1155					
GAC	AGT	GAA	ACA	GTA	GAA	CCT	ATT	ATC	TCA	CAG	TTG	GTG	ACA	GTG	CTT	3857
Asp	Ser	Glu	Thr	Val	Glu	Pro	Ile	Ile	Ser	Gln	Leu	Val	Thr	Val	Leu	
1160					1165					1170					1175	
TTG	AAA	GGT	TGC	CAA	GAT	GCA	AAC	TCT	CAA	GCT	CGG	TTG	CTC	TGT	GGG	3905
Leu	Lys	Gly	Cys	Gln	Asp	Ala	Asn	Ser	Gln	Ala	Arg	Leu	Leu	Cys	Gly	
				1180					1185					1190		
GAA	TGT	TTA	GGG	GAA	TTG	GGG	GCG	ATA	GAT	CCA	GGT	CGA	TTA	GAT	TTC	3953
Glu	Cys	Leu	Gly	Glu	Leu	Gly	Ala	Ile	Asp	Pro	Gly	Arg	Leu	Asp	Phe	
			1195					1200					1205			
TCA	ACA	ACT	GAA	ACT	CAA	GGA	AAA	GAT	TTT	ACA	TTT	GTG	ACT	GGA	GTA	4001
Ser	Thr	Thr	Glu	Thr	Gln	Gly	Lys	Asp	Phe	Thr	Phe	Val	Thr	Gly	Val	
			1210				1215					1220				
GAA	GAT	TCA	AGC	TTT	GCC	TAT	GGA	TTA	TTG	ATG	GAG	CTA	ACA	AGA	GCT	4049
Glu	Asp	Ser	Ser	Phe	Ala	Tyr	Gly	Leu	Leu	Met	Glu	Leu	Thr	Arg	Ala	
	1225					1230					1235					
TAC	CTT	GCG	TAT	GCT	GAT	AAT	AGC	CGA	GCT	CAA	GAT	TCA	GCT	GCC	TAT	4097
Tyr	Leu	Ala	Tyr	Ala	Asp	Asn	Ser	Arg	Ala	Gln	Asp	Ser	Ala	Ala	Tyr	
1240					1245					1250					1255	
GCC	ATT	CAG	GAG	TTG	CTT	TCT	ATT	TAT	GAC	TGT	AGA	GAG	ATG	GAG	ACC	4145
Ala	Ile	Gln	Glu	Leu	Leu	Ser	Ile	Tyr	Asp	Cys	Arg	Glu	Met	Glu	Thr	
				1260					1265					1270		
AAC	GGC	CCA	GGT	CAC	CAA	TTG	TGG	AGG	AGA	TTT	CCT	GAG	CAT	GTT	CGG	4193
Asn	Gly	Pro	Gly	His	Gln	Leu	Trp	Arg	Arg	Phe	Pro	Glu	His	Val	Arg	
			1275					1280					1285			
GAA	ATA	CTA	GAA	CCT	CAT	CTA	AAT	ACC	AGA	TAC	AAG	AGT	TCT	CAG	AAG	4241
Glu	Ile	Leu	Glu	Pro	His	Leu	Asn	Thr	Arg	Tyr	Lys	Ser	Ser	Gln	Lys	
			1290				1295					1300				
TCA	ACC	GAT	TGG	TCT	GGA	GTA	AAG	AAG	CCA	ATT	TAC	TTA	AGT	AAA	TTG	4289
Ser	Thr	Asp	Trp	Ser	Gly	Val	Lys	Lys	Pro	Ile	Tyr	Leu	Ser	Lys	Leu	
	1305					1310					1315					

-102-

GGT AGT AAC TTT GCA GAA TGG TCA GCA TCT TGG GCA GGT TAT CTT ATT Gly Ser Asn Phe Ala Glu Trp Ser Ala Ser Trp Ala Gly Tyr Leu Ile 1320 1325 1330 1335	4337
ACA AAG GTT CGA CAT GAT CTT GCC AGT AAA ATT TTC ACC TGC TGT AGC Thr Lys Val Arg His Asp Leu Ala Ser Lys Ile Phe Thr Cys Cys Ser 1340 1345 1350	4385
ATT ATG ATG AAG CAT GAT TTC AAA GTG ACC ATC TAT CTT CTT CCA CAT Ile Met Met Lys His Asp Phe Lys Val Thr Ile Tyr Leu Leu Pro His 1355 1360 1365	4433
ATT CTG GTG TAT GTC TTA CTG GGT TGT AAT CAA GAA GAT CAG CAG GAG Ile Leu Val Tyr Val Leu Leu Gly Cys Asn Gln Glu Asp Gln Gln Glu 1370 1375 1380	4481
GTT TAT GCA GAA ATT ATG GCA GTT CTA AAG CAT GAC GAT CAG CAT ACC Val Tyr Ala Glu Ile Met Ala Val Leu Lys His Asp Asp Gln His Thr 1385 1390 1395	4529
ATA AAT ACC CAA GAC ATT GCA TCT GAT CTG TGT CAA CTC AGT ACA CAG Ile Asn Thr Gln Asp Ile Ala Ser Asp Leu Cys Gln Leu Ser Thr Gln 1400 1405 1410 1415	4577
ACT GTG TTC TCC ATG CTT GAC CAT CTC ACA CAG TGG GCA AGG CAC AAA Thr Val Phe Ser Met Leu Asp His Leu Thr Gln Trp Ala Arg His Lys 1420 1425 1430	4625
TTT CAG GCA CTG AAA GCT GAG AAA TGT CCA CAC AGC AAA TCA AAC AGA Phe Gln Ala Leu Lys Ala Glu Lys Cys Pro His Ser Lys Ser Asn Arg 1435 1440 1445	4673
AAT AAG GTA GAC TCA ATG GTA TCT ACT GTG GAT TAT GAA GAC TAT CAG Asn Lys Val Asp Ser Met Val Ser Thr Val Asp Tyr Glu Asp Tyr Gln 1450 1455 1460	4721
AGT GTA ACC CGT TTT CTA GAC CTC ATA CCC CAG GAT ACT CTG GCA GTA Ser Val Thr Arg Phe Leu Asp Leu Ile Pro Gln Asp Thr Leu Ala Val 1465 1470 1475	4769
GCT TCC TTT CGC TCC AAA GCA TAC ACA CGA GCT GTA ATG CAC TTT GAA Ala Ser Phe Arg Ser Lys Ala Tyr Thr Arg Ala Val Met His Phe Glu 1480 1485 1490 1495	4817
TCA TTT ATT ACA GAA AAG AAG CAA AAT ATT CAG GAA CAT CTT GGA TTT Ser Phe Ile Thr Glu Lys Lys Gln Asn Ile Gln Glu His Leu Gly Phe 1500 1505 1510	4865
TTA CAG AAA TTG TAT GCT GCT ATG CAT GAA CCT GAT GGA GTG GCC GGA Leu Gln Lys Leu Tyr Ala Ala Met His Glu Pro Asp Gly Val Ala Gly 1515 1520 1525	4913
GTC AGT GCA ATT AGA AAG GCA GAA CCA TCT CTA AAA GAA CAG ATC CTT Val Ser Ala Ile Arg Lys Ala Glu Pro Ser Leu Lys Glu Gln Ile Leu 1530 1535 1540	4961
GAA CAT GAA AGC CTT GGC TTG CTG AGG GAT GCC ACT GCT TGT TAT GAC Glu His Glu Ser Leu Gly Leu Leu Arg Asp Ala Thr Ala Cys Tyr Asp 1545 1550 1555	5009
AGG GCT ATT CAG CTA GAA CCA GAC CAG ATC ATT CAT TAC CAT GGT GTA Arg Ala Ile Gln Leu Glu Pro Asp Gln Ile Ile His Tyr His Gly Val 1560 1565 1570 1575	5057
GTA AAG TCC ATG TTA GGT CTT GGT CAG CTG TCT ACT GTT ATC ACT CAG Val Lys Ser Met Leu Gly Leu Gly Gln Leu Ser Thr Val Ile Thr Gln 1580 1585 1590	5105

-103-

GTG AAT GGA GTG CAT GCT AAC AGG TCC GAG TGG ACA GAT GAA TTA AAC	5153
Val Asn Gly Val His Ala Asn Arg Ser Glu Trp Thr Asp Glu Leu Asn	
1595 1600 1605	
ACG TAC AGA GTG GAA GCA GCT TGG AAA TTG TCA CAG TGG GAT TTG GTG	5201
Thr Tyr Arg Val Glu Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val	
1610 1615 1620	
GAA AAC TAT TTG GCA GCA GAT GGA AAA TCT ACA ACA TGG AGT GTC AGA	5249
Glu Asn Tyr Leu Ala Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg	
1625 1630 1635	
CTG GGA CAG CTA TTA TTA TCA GCC AAA AAA AGA GAT ATC ACA GCT TTT	5297
Leu Gly Gln Leu Leu Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe	
1640 1645 1650 1655	
TAT GAC TCA CTG AAA CTA GTG AGA GCA GAA CAA ATT GTA CCT CTT TCA	5345
Tyr Asp Ser Leu Lys Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser	
1660 1665 1670	
GCT GCA AGC TTT GAA AGA GGC TCC TAC CAA CGA GGA TAT GAA TAT ATT	5393
Ala Ala Ser Phe Glu Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile	
1675 1680 1685	
GTG AGA TTG CAC ATG TTA TGT GAG TTG GAG CAT AGC ATC AAA CCA CTT	5441
Val Arg Leu His Met Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu	
1690 1695 1700	
TTC CAG CAT TCT CCA GGT GAC AGT TCT CAA GAA GAT TCT CTA AAC TGG	5489
Phe Gln His Ser Pro Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp	
1705 1710 1715	
GTA GCT CGA CTA GAA ATG ACC CAG AAT TCC TAC AGA GCC AAG GAG CCT	5537
Val Ala Arg Leu Glu Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro	
1720 1725 1730 1735	
ATC CTG GCT CTC CGG AGG GCT TTA CTA AGC CTC AAC AAA AGA CCA GAT	5585
Ile Leu Ala Leu Arg Arg Ala Leu Leu Ser Leu Asn Lys Arg Pro Asp	
1740 1745 1750	
TAC AAT GAA ATG GTT GGA GAA TGC TGG CTG CAG AGT GCC AGG GTA GCT	5633
Tyr Asn Glu Met Val Gly Glu Cys Trp Leu Gln Ser Ala Arg Val Ala	
1755 1760 1765	
AGA AAG GCT GGT CAC CAC CAG ACA GCC TAC AAT GCT CTC CTT AAT GCA	5681
Arg Lys Ala Gly His His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala	
1770 1775 1780	
GGG GAA TCA CGA CTC GCT GAA CTG TAC GTG GAA AGG GCA AAG TGG CTC	5729
Gly Glu Ser Arg Leu Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu	
1785 1790 1795	
TGG TCC AAG GGT GAT GTT CAC CAG GCA CTA ATT GTT CTT CAA AAA GGT	5777
Trp Ser Lys Gly Asp Val His Gln Ala Leu Ile Val Leu Gln Lys Gly	
1800 1805 1810 1815	
GTT GAA TTA TGT TTT CCT GAA AAT GAA ACC CCA CCT GAG GGT AAG AAC	5825
Val Glu Leu Cys Phe Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn	
1820 1825 1830	
ATG TTA ATC CAT GGT CGA GCT ATG CTA CTA GTG GGC CGA TTT ATG GAA	5873
Met Leu Ile His Gly Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu	
1835 1840 1845	
GAA ACA GCT AAC TTT GAA AGC AAT GCA ATT ATG AAA AAA TAT AAG GAT	5921
Glu Thr Ala Asn Phe Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp	
1850 1855 1860	

-104-

GTG ACC GCG TGC CTG CCA GAA TGG GAG GAT GGG CAT TTT TAC CTT GCC Val Thr Ala Cys Leu Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala 1865 1870 1875	5969
AAG TAC TAT GAC AAA TTG ATG CCC ATG GTC ACA GAC AAC AAA ATG GAA Lys Tyr Tyr Asp Lys Leu Met Pro Met Val Thr Asp Asn Lys Met Glu 1880 1885 1890 1895	6017
AAG CAA GGT GAT CTC ATC CGG TAT ATA GTT CTT CAT TTT GGC AGA TCT Lys Gln Gly Asp Leu Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser 1900 1905 1910	6065
CTA CAA TAT GGA AAT CAG TTC ATA TAT CAG TCA ATG CCA CGA ATG TTA Leu Gln Tyr Gly Asn Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu 1915 1920 1925	6113
ACT CTA TGG CTT GAT TAT GGT ACA AAG GCA TAT GAA TGG GAA AAA GCT Thr Leu Trp Leu Asp Tyr Gly Thr Lys Ala Tyr Glu Trp Glu Lys Ala 1930 1935 1940	6161
GGC CGC TCC GAT CGT GTA CAA ATG AGG AAT GAT TTG GGT AAA ATA AAC Gly Arg Ser Asp Arg Val Gln Met Arg Asn Asp Leu Gly Lys Ile Asn 1945 1950 1955	6209
AAG GTT ATC ACA GAG CAT ACA AAC TAT TTA GCT CCA TAT CAA TTT TTG Lys Val Ile Thr Glu His Thr Asn Tyr Leu Ala Pro Tyr Gln Phe Leu 1960 1965 1970 1975	6257
ACT GCT TTT TCA CAA TTG ATC TCT CGA ATT TGT CAT TCT CAC GAT GAA Thr Ala Phe Ser Gln Leu Ile Ser Arg Ile Cys His Ser His Asp Glu 1980 1985 1990	6305
GTT TTT GTT GTG CTT GAT GGA AAT AAT AGC CAA GTA TTT CTA GCC TAT Val Phe Val Val Leu Asp Gly Asn Asn Ser Gln Val Phe Leu Ala Tyr 1995 2000 2005	6353
CCT CAA CAA GCA ATG TGG ATG ATG ACA GCT GTG TCA AAG TCA TCT TAT Pro Gln Gln Ala Met Trp Met Met Thr Ala Val Ser Lys Ser Ser Tyr 2010 2015 2020	6401
CCC ATG CGT GTG AAC AGA TGC AAG GAA ATC CTC AAT AAA GCT ATT CAT Pro Met Arg Val Asn Arg Cys Lys Glu Ile Leu Asn Lys Ala Ile His 2025 2030 2035	6449
ATG AAA AAA TCC TTA GAG AAG TTT GTT GGA GAT GCA ACT CGC CTA ACA Met Lys Lys Ser Leu Glu Lys Phe Val Gly Asp Ala Thr Arg Leu Thr 2040 2045 2050 2055	6497
GAT AAG CTT CTA GAA TTG TGC AAT AAA CCG GTG GAA ATT CTT GCT TCT Asp Lys Leu Leu Glu Leu Cys Asn Lys Pro Val Glu Ile Leu Ala Ser 2060 2065 2070	6545
CTT CAG AAA CCA AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG TTC Leu Gln Lys Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe 2075 2080 2085	6593
TAC ATC ATG ATG TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT AGA Tyr Ile Met Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg 2090 2095 2100	6641
CTA ATG GAA TTC AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT GCA Leu Met Glu Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala 2105 2110 2115	6689
GAG TCT CGT AGA AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT CCA Glu Ser Arg Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro 2120 2125 2130 2135	6737

-105-

CTA AAT GAT GAA TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT GGT Leu Asn Asp Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly 2140 2145 2150	6785
TTG AGA CCT ATT CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT ATG Leu Arg Pro Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met 2155 2160 2165	6833
ACA GGA AAA GAA CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT TTA Thr Gly Lys Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu 2170 2175 2180	6881
TCT GAA AAA CTC AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT CCT Ser Glu Lys Leu Lys Val Phe Arg Glu Phe Leu Pro Arg His Pro 2185 2190 2195	6929
CCT ATT TTT CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA TCA Pro Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser 2200 2205 2210 2215	6977
TGG TAC AGT AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser 2220 2225 2230	7025
ATG GTT GGT TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile 2235 2240 2245	7073
CTC TTT GAT TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys 2250 2255 2260	7121
CTT TTC AAT AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe 2265 2270 2275	7169
CGC CTG ACT CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu 2280 2285 2290 2295	7217
GGT CTT TTT CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp 2300 2305 2310	7265
CAG CGA GAG CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro 2315 2320 2325	7313
CTT GTG GAA TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu 2330 2335 2340	7361
AAT GAA ACT GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu 2345 2350 2355	7409
GAC ATT GAG CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val 2360 2365 2370 2375	7457
ACA GGA CTG CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln 2380 2385 2390	7505
GAA GCT ACT GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr 2395 2400 2405	7553

-106-

CCA TAT ATG TGAAATGAAA TTATGTAAAA GAATATGTGA ATAATCTAAA  
 Pro Tyr Met  
 2410

7602

AGTAAAAAAA AAAAAAAAAA AA

7624

## (2) INFORMATION FOR SEQ ID NO:24:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2410 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

Met	Gly	His	Ala	Val	Glu	Trp	Pro	Val	Val	Met	Ser	Arg	Phe	Leu	Ser	1	5	10	15
Gln	Leu	Asp	Glu	His	Met	Gly	Tyr	Leu	Gln	Ser	Ala	Pro	Leu	Gln	Leu	20	25	30	
Met	Ser	Met	Gln	Asn	Leu	Glu	Phe	Ile	Glu	Val	Thr	Leu	Leu	Met	Val	35	40	45	
Leu	Thr	Arg	Ile	Ile	Ala	Ile	Val	Phe	Phe	Arg	Arg	Gln	Glu	Leu	Leu	50	55	60	
Leu	Trp	Gln	Ile	Gly	Cys	Val	Leu	Leu	Glu	Tyr	Gly	Ser	Pro	Lys	Ile	65	70	75	80
Lys	Ser	Leu	Ala	Ile	Ser	Phe	Leu	Thr	Glu	Leu	Phe	Gln	Leu	Gly	Gly	85	90	95	
Leu	Pro	Ala	Gln	Pro	Ala	Ser	Thr	Phe	Phe	Ser	Ser	Phe	Leu	Glu	Leu	100	105	110	
Leu	Lys	His	Leu	Val	Glu	Met	Asp	Thr	Asp	Gln	Leu	Lys	Leu	Tyr	Glu	115	120	125	
Glu	Pro	Leu	Ser	Lys	Leu	Ile	Lys	Thr	Leu	Phe	Pro	Phe	Glu	Ala	Glu	130	135	140	
Ala	Tyr	Arg	Asn	Ile	Glu	Pro	Val	Tyr	Leu	Asn	Met	Leu	Leu	Glu	Lys	145	150	155	160
Leu	Cys	Val	Met	Phe	Glu	Asp	Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	165	170	175	
Leu	Leu	Lys	Ala	Ala	Leu	Cys	His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	180	185	190	
Val	Pro	Ala	Gly	Tyr	Glu	Ser	Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	195	200	205	
Arg	Asn	Ile	Cys	Lys	Ala	Leu	Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	210	215	220	
Ala	Glu	Tyr	Leu	Leu	Gly	Pro	Leu	Tyr	Ala	Ala	Leu	Lys	Met	Glu	Ser	225	230	235	240
Met	Glu	Ile	Ile	Glu	Glu	Ile	Gln	Cys	Gln	Thr	Gln	Gln	Glu	Asn	Leu	245	250	255	



-107-

Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg Arg Leu Ser Ser  
 260 265 270  
 Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr Glu Glu Ile Lys  
 275 280 285  
 His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser Ala Leu Lys Gln  
 290 295 300  
 Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser Gly Leu Lys Asn  
 305 310 315 320  
 Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val Leu Gln Leu Thr  
 325 330 335  
 Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met Asn Cys Arg Thr  
 340 345 350  
 Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro Ser Val Val Ile  
 355 360 365  
 Thr Trp Met Ser Leu Asp Phe Tyr Thr Lys Val Leu Lys Ser Cys Arg  
 370 375 380  
 Ser Leu Leu Glu Ser Val Gln Lys Leu Asp Leu Glu Ala Thr Ile Asp  
 385 390 395 400  
 Lys Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met Gln Val Asn Ser  
 405 410 415  
 Ser Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly Met Leu Ser Leu  
 420 425 430  
 Pro Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu Lys Leu Thr Thr  
 435 440 445  
 Phe Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile Ser Asp Ser Tyr  
 450 455 460  
 Ser Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu Thr Leu Phe Pro  
 465 470 475 480  
 Arg Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr Asn Trp Ala Leu  
 485 490 495  
 Gln Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe  
 500 505 510  
 Ile Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu  
 515 520 525  
 Ile Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala  
 530 535 540  
 Ser Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu  
 545 550 555 560  
 Thr Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu  
 565 570 575  
 Phe Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser  
 580 585 590  
 Gln Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe Leu Leu Lys Lys  
 595 600 605

-108-

Lys Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp Asn Leu His His  
 610 615 620  
 Leu Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr Asp Val Lys Ala  
 625 630 635 640  
 Val Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro Asp Lys Asp Val  
 645 650 655  
 Arg Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu Glu Ser Leu Asp  
 660 665 670  
 Ser Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu Arg Met Lys Glu  
 675 680 685  
 Ala Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu Leu Lys Asp Thr  
 690 695 700  
 Leu Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala Lys Gly Asp Leu  
 705 710 715 720  
 Val Pro Phe Ala Leu Leu His Leu Leu His Cys Leu Leu Ser Lys Ser  
 725 730 735  
 Ala Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg Ala Leu Val Ala  
 740 745 750  
 Ala Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln Tyr Lys Lys Pro  
 755 760 765  
 Ile Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser Gln Met Thr Ala  
 770 775 780  
 Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val  
 785 790 795 800  
 Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn  
 805 810 815  
 Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln  
 820 825 830  
 Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala  
 835 840 845  
 Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile  
 850 855 860  
 Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys  
 865 870 875 880  
 Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu Lys Asn Glu Thr  
 885 890 895  
 Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe Gln Gly Leu His  
 900 905 910  
 Asn Glu Leu Leu Leu Arg Ile Gly Glu His Tyr Gln Gln Val Phe Asn  
 915 920 925  
 Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp Asp Pro Tyr Gln  
 930 935 940  
 Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln  
 945 950 955 960

-109-

Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser  
 965 970 975  
 Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met  
 980 985 990  
 Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val  
 995 1000 1005  
 Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe  
 1010 1015 1020  
 Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp  
 1025 1030 1035 1040  
 His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu  
 1045 1050 1055  
 Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr  
 1060 1065 1070  
 Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile  
 1075 1080 1085  
 Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu  
 1090 1095 1100  
 Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr  
 1105 1110 1115 1120  
 Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg  
 1125 1130 1135  
 Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu  
 1140 1145 1150  
 Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile  
 1155 1160 1165  
 Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser  
 1170 1175 1180  
 Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile  
 1185 1190 1195 1200  
 Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp  
 1205 1210 1215  
 Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu  
 1220 1225 1230  
 Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg  
 1235 1240 1245  
 Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr  
 1250 1255 1260  
 Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg  
 1265 1270 1275 1280  
 Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr  
 1285 1290 1295  
 Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys  
 1300 1305 1310

-110-

Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala  
 1315 1320 1325  
 Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser  
 1330 1335 1340  
 Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val  
 1345 1350 1355 1360  
 Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys  
 1365 1370 1375  
 Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu  
 1380 1385 1390  
 Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp  
 1395 1400 1405  
 Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu  
 1410 1415 1420  
 Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys  
 1425 1430 1435 1440  
 Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr  
 1445 1450 1455  
 Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile  
 1460 1465 1470  
 Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr  
 1475 1480 1485  
 Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn  
 1490 1495 1500  
 Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His  
 1505 1510 1515 1520  
 Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg Lys Ala Glu Pro  
 1525 1530 1535  
 Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg  
 1540 1545 1550  
 Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln  
 1555 1560 1565  
 Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln  
 1570 1575 1580  
 Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser  
 1585 1590 1595 1600  
 Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys  
 1605 1610 1615  
 Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys  
 1620 1625 1630  
 Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys  
 1635 1640 1645  
 Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala  
 1650 1655 1660

-111-

Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr  
 1665 1670 1675 1680  
 Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu  
 1685 1690 1695  
 Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser  
 1700 1705 1710  
 Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn  
 1715 1720 1725  
 Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu  
 1730 1735 1740  
 Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp  
 1745 1750 1755 1760  
 Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala  
 1765 1770 1775  
 Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr  
 1780 1785 1790  
 Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala  
 1795 1800 1805  
 Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu  
 1810 1815 1820  
 Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu  
 1825 1830 1835 1840  
 Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala  
 1845 1850 1855  
 Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu  
 1860 1865 1870  
 Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met  
 1875 1880 1885  
 Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile  
 1890 1895 1900  
 Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr  
 1905 1910 1915 1920  
 Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys  
 1925 1930 1935  
 Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg  
 1940 1945 1950  
 Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr  
 1955 1960 1965  
 Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg  
 1970 1975 1980  
 Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn  
 1985 1990 1995 2000  
 Ser Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr  
 2005 2010 2015

-112-

Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu  
2020 2025 2030

Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val  
2035 2040 2045

Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys  
2050 2055 2060

Pro Val Glu Ile Leu Ala Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu  
2065 2070 2075 2080

Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met Met Cys Lys Pro Lys Asp  
2085 2090 2095

Asp Leu Arg Lys Asp Cys Arg Leu Met Glu Phe Asn Ser Leu Ile Asn  
2100 2105 2110

Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg Arg Arg Glu Leu His Ile  
2115 2120 2125

Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu  
2130 2135 2140

Trp Val Asn Asn Thr Ala Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr  
2145 2150 2155 2160

Lys Glu Lys Gly Val Tyr Met Thr Gly Lys Glu Leu Arg Gln Cys Met  
2165 2170 2175

Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys Leu Lys Val Phe Arg Glu  
2180 2185 2190

Phe Leu Leu Pro Arg His Pro Pro Ile Phe His Glu Trp Phe Leu Arg  
2195 2200 2205

Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys  
2210 2215 2220

Arg Ser Thr Ala Val Met Ser Met Val Gly Tyr Ile Leu Gly Leu Gly  
2225 2230 2235 2240

Asp Arg His Gly Glu Asn Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys  
2245 2250 2255

Val His Val Asp Phe Asn Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu  
2260 2265 2270

Val Pro Glu Ile Val Pro Phe Arg Leu Thr His Asn Met Val Asn Gly  
2275 2280 2285

Met Gly Pro Met Gly Thr Glu Gly Leu Phe Arg Arg Ala Cys Glu Val  
2290 2295 2300

Thr Met Arg Leu Met Arg Asp Gln Arg Glu Pro Leu Met Ser Val Leu  
2305 2310 2315 2320

Lys Thr Phe Leu His Asp Pro Leu Val Glu Trp Ser Lys Pro Val Lys  
2325 2330 2335

Gly His Ser Lys Ala Pro Leu Asn Glu Thr Gly Glu Val Val Asn Glu  
2340 2345 2350

Lys Ala Lys Thr His Val Leu Asp Ile Glu Gln Arg Leu Gln Gly Val  
2355 2360 2365

-113-

Ile Lys Thr Arg Asn Arg Val Thr Gly Leu Pro Leu Ser Ile Glu Gly  
 2370 2375 2380

His Val His Tyr Leu Ile Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys  
 2385 2390 2395 2400

Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2405 2410

## (2) INFORMATION FOR SEQ ID NO:25:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7502 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..7440

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

ATG GGT CAT GCT GTG GAA TGG CCA GTG GTC ATG AGC CGA TTT TTA AGT	48
Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser	
1 5 10 15	
CAA TTA GAT GAA CAC ATG GGA TAT TTA CAA TCA GCT CCT TTG CAG TTG	96
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu	
20 25 30	
ATG AGT ATG CAA AAT TTA GAA TTT ATT GAA GTC ACT TTA TTA ATG GTT	144
Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr Leu Leu Met Val	
35 40 45	
CTT ACT CGT ATT ATT GCA ATT GTG TTT TTT AGA AGG CAA GAA CTC TTA	192
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Gln Glu Leu Leu	
50 55 60	
CTT TGG CAG ATA GGT TGT GTT CTG CTA GAG TAT GGT AGT CCA AAA ATT	240
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile	
65 70 75 80	
AAA TCC CTA GCA ATT AGC TTT TTA ACA GAA CTT TTT CAG CTT GGA GGA	288
Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly	
85 90 95	
CTA CCA GCA CAA CCA GCT AGC ACT TTT TTC AGC TCA TTT TTG GAA TTA	336
Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu	
100 105 110	
TTA AAA CAC CTT GTA GAA ATG GAT ACT GAC CAA TTG AAA CTC TAT GAA	384
Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu	
115 120 125	
GAG CCA TTA TCA AAG CTG ATA AAG ACA CTA TTT CCC TTT GAA GCA GAA	432
Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu	
130 135 140	
GCT TAT AGA AAT ATT GAA CCT GTC TAT TTA AAT ATG CTG CTG GAA AAA	480
Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys	
145 150 155 160	

-114-

CTC	TGT	GTC	ATG	TTT	GAA	GAC	GGT	GTG	CTC	ATG	CGG	CTT	AAG	TCT	GAT	528
Leu	Cys	Val	Met	Phe	Glu	Asp	Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	
				165					170					175		
TTG	CTA	AAA	GCA	GCT	TTG	TGC	CAT	TTA	CTG	CAG	TAT	TTC	CTT	AAA	TTT	576
Leu	Leu	Lys	Ala	Ala	Leu	Cys	His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	
			180					185					190			
GTG	CCA	GCT	GGG	TAT	GAA	TCT	GCT	TTA	CAA	GTC	AGG	AAG	GTC	TAT	GTG	624
Val	Pro	Ala	Gly	Tyr	Glu	Ser	Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	
		195					200					205				
AGA	AAT	ATT	TGT	AAA	GCT	CTT	TTG	GAT	GTG	CTT	GGA	ATT	GAG	GTA	GAT	672
Arg	Asn	Ile	Cys	Lys	Ala	Leu	Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	
	210					215					220					
GCA	GAG	TAC	TTG	TTG	GGC	CCA	CTT	TAT	GCA	GCT	TTG	AAA	ATG	GAA	AGT	720
Ala	Glu	Tyr	Leu	Leu	Gly	Pro	Leu	Tyr	Ala	Ala	Leu	Lys	Met	Glu	Ser	
225					230				235						240	
ATG	GAA	ATC	ATT	GAG	GAG	ATT	CAA	TGC	CAA	ACT	CAA	CAG	GAA	AAC	CTC	768
Met	Glu	Ile	Ile	Glu	Glu	Ile	Gln	Cys	Gln	Thr	Gln	Gln	Glu	Asn	Leu	
				245				250						255		
AGC	AGT	AAT	AGT	GAT	GGA	ATA	TCA	CCC	AAA	AGG	CGT	CGT	CTC	AGC	TCG	816
Ser	Ser	Asn	Ser	Asp	Gly	Ile	Ser	Pro	Lys	Arg	Arg	Arg	Leu	Ser	Ser	
			260					265					270			
TCT	CTA	AAC	CCT	TCT	AAA	AGA	GCA	CCA	AAA	CAG	ACT	GAG	GAA	ATT	AAA	864
Ser	Leu	Asn	Pro	Ser	Lys	Arg	Ala	Pro	Lys	Gln	Thr	Glu	Glu	Ile	Lys	
		275					280					285				
CAT	GTG	GAC	ATG	AAC	CAA	AAG	AGC	ATA	TTA	TGG	AGT	GCA	CTG	AAA	CAG	912
His	Val	Asp	Met	Asn	Gln	Lys	Ser	Ile	Leu	Trp	Ser	Ala	Leu	Lys	Gln	
	290					295					300					
AAA	GCT	GAA	TCC	CTT	CAG	ATT	TCC	CTT	GAA	TAC	AGT	GGC	CTA	AAG	AAT	960
Lys	Ala	Glu	Ser	Leu	Gln	Ile	Ser	Leu	Glu	Tyr	Ser	Gly	Leu	Lys	Asn	
305					310				315						320	
CCT	GTT	ATT	GAG	ATG	TTA	GAA	GGA	ATT	GCT	GTT	GTC	TTA	CAA	CTG	ACT	1008
Pro	Val	Ile	Glu	Met	Leu	Glu	Gly	Ile	Ala	Val	Val	Leu	Gln	Leu	Thr	
				325					330					335		
GCT	CTG	TGT	ACT	GTT	CAT	TGT	TCT	CAT	CAA	AAC	ATG	AAC	TGC	CGT	ACT	1056
Ala	Leu	Cys	Thr	Val	His	Cys	Ser	His	Gln	Asn	Met	Asn	Cys	Arg	Thr	
			340				345						350			
TTC	AAG	GAC	TGT	CAA	CAT	AAA	TCC	AAG	AAG	AAA	CCT	TCT	GTA	GTG	ATA	1104
Phe	Lys	Asp	Cys	Gln	His	Lys	Ser	Lys	Lys	Lys	Pro	Ser	Val	Val	Ile	
		355					360					365				
ACT	TGG	ATG	TCA	TTG	GAT	TTT	TAC	ACA	AAA	GTG	CTT	AAG	AGC	TGT	AGA	1152
Thr	Trp	Met	Ser	Leu	Asp	Phe	Tyr	Thr	Lys	Val	Leu	Lys	Ser	Cys	Arg	
	370					375					380					
AGT	TTG	TTA	GAA	TCT	GTT	CAG	AAA	CTG	GAC	CTG	GAG	GCA	ACC	ATT	GAT	1200
Ser	Leu	Leu	Glu	Ser	Val	Gln	Lys	Leu	Asp	Leu	Glu	Ala	Thr	Ile	Asp	
385					390				395						400	
AAG	GTG	GTG	AAA	ATT	TAT	GAT	GCT	TTG	ATT	TAT	ATG	CAA	GTA	AAC	AGT	1248
Lys	Val	Val	Lys	Ile	Tyr	Asp	Ala	Leu	Ile	Tyr	Met	Gln	Val	Asn	Ser	
			405						410					415		
TCA	TTT	GAA	GAT	CAT	ATC	CTG	GAA	GAT	TTA	TGT	GGA	ATG	CTC	TCA	CTT	1296
Ser	Phe	Glu	Asp	His	Ile	Leu	Glu	Asp	Leu	Cys	Gly	Met	Leu	Ser	Leu	
			420				425						430			



-115-

CCA	TGG	ATT	TAT	TCC	CAT	TCT	GAT	GAT	GGC	TGT	TTA	AAG	TTG	ACC	ACA	1344
Pro	Trp	Ile	Tyr	Ser	His	Ser	Asp	Asp	Gly	Cys	Leu	Lys	Leu	Thr	Thr	
	435						440					445				
TTT	GCC	GCT	AAT	CTT	CTA	ACA	TTA	AGC	TGT	AGG	ATT	TCA	GAT	AGC	TAT	1392
Phe	Ala	Ala	Asn	Leu	Leu	Thr	Leu	Ser	Cys	Arg	Ile	Ser	Asp	Ser	Tyr	
	450					455					460					
TCA	CCA	CAG	GCA	CAA	TCA	CGA	TGT	GTG	TTT	CTT	CTG	ACT	CTG	TTT	CCA	1440
Ser	Pro	Gln	Ala	Gln	Ser	Arg	Cys	Val	Phe	Leu	Thr	Leu	Thr	Phe	Pro	
465					470					475					480	
AGA	AGA	ATA	TTC	CTT	GAG	TGG	AGA	ACA	GCA	GTT	TAC	AAC	TGG	GCC	CTG	1488
Arg	Arg	Ile	Phe	Leu	Glu	Trp	Arg	Thr	Ala	Val	Tyr	Asn	Trp	Ala	Leu	
				485					490					495		
CAG	AGC	TCC	CAT	GAA	GTA	ATC	CGG	GCT	AGT	TGT	GTT	AGT	GGA	TTT	TTT	1536
Gln	Ser	Ser	His	Glu	Val	Ile	Arg	Ala	Ser	Cys	Val	Ser	Gly	Phe	Phe	
			500					505					510			
ATC	TTA	TTG	CAG	CAG	CAG	AAT	TCT	TGT	AAC	AGA	GTT	CCC	AAG	ATT	CTT	1584
Ile	Leu	Leu	Gln	Gln	Gln	Asn	Ser	Cys	Asn	Arg	Val	Pro	Lys	Ile	Leu	
	515						520					525				
ATA	GAT	AAA	GTC	AAA	GAT	GAT	TCT	GAC	ATT	GTC	AAG	AAA	GAA	TTT	GCT	1632
Ile	Asp	Lys	Val	Lys	Asp	Asp	Ser	Asp	Ile	Val	Lys	Lys	Glu	Phe	Ala	
	530					535					540					
TCT	ATA	CTT	GGT	CAA	CTT	GTC	TGT	ACT	CTT	CAC	GGC	ATG	TTT	TAT	CTG	1680
Ser	Ile	Leu	Gly	Gln	Leu	Val	Cys	Thr	Leu	His	Gly	Met	Phe	Tyr	Leu	
545					550					555					560	
ACA	AGT	TCT	TTA	ACA	GAA	CCT	TTC	TCT	GAA	CAC	GGA	CAT	GTG	GAC	CTC	1728
Thr	Ser	Ser	Leu	Thr	Glu	Pro	Phe	Ser	Glu	His	Gly	His	Val	Asp	Leu	
				565					570					575		
TTC	TGT	AGG	AAC	TTG	AAA	GCC	ACT	TCT	CAA	CAT	GAA	TGT	TCA	TCT	TCT	1776
Phe	Cys	Arg	Asn	Leu	Lys	Ala	Thr	Ser	Gln	His	Glu	Cys	Ser	Ser	Ser	
			580					585					590			
CAA	CTA	AAA	GCT	TCT	GTC	TGC	AAG	CCA	TTC	CTT	TTC	CTA	CTG	AAA	AAA	1824
Gln	Leu	Lys	Ala	Ser	Val	Cys	Lys	Pro	Phe	Leu	Phe	Leu	Leu	Lys	Lys	
	595						600					605				
AAA	ATA	CCT	AGT	CCA	GTA	AAA	CTT	GCT	TTC	ATA	GAT	AAT	CTA	CAT	CAT	1872
Lys	Ile	Pro	Ser	Pro	Val	Lys	Leu	Ala	Phe	Ile	Asp	Asn	Leu	His	His	
	610					615					620					
CTT	TGT	AAG	CAT	CTT	GAT	TTT	AGA	GAA	GAT	GAA	ACA	GAT	GTA	AAA	GCA	1920
Leu	Cys	Lys	His	Leu	Asp	Phe	Arg	Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala	
625					630					635					640	
GTT	CTT	GGA	ACT	TTA	TTA	AAT	TTA	ATG	GAA	GAT	CCA	GAC	AAA	GAT	GTT	1968
Val	Leu	Gly	Thr	Leu	Leu	Asn	Leu	Met	Glu	Asp	Pro	Asp	Lys	Asp	Val	
				645					650					655		
AGA	GTG	GCT	TTT	AGT	GGA	AAT	ATC	AAG	CAC	ATA	TTG	GAA	TCC	TTG	GAC	2016
Arg	Val	Ala	Phe	Ser	Gly	Asn	Ile	Lys	His	Ile	Leu	Glu	Ser	Leu	Asp	
			660					665					670			
TCT	GAA	GAT	GGA	TTT	ATA	AAG	GAG	CTT	TTT	GTC	TTA	AGA	ATG	AAG	GAA	2064
Ser	Glu	Asp	Gly	Phe	Ile	Lys	Glu	Leu	Phe	Val	Leu	Arg	Met	Lys	Glu	
		675					680					685				
GCA	TAT	ACA	CAT	GCC	CAA	ATA	TCA	AGA	AAT	AAT	GAG	CTG	AAG	GAT	ACC	2112
Ala	Tyr	Thr	His	Ala	Gln	Ile	Ser	Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr	
	690					695					700					

-116-

TTG	ATT	CTT	ACA	ACA	GGG	GAT	ATT	GGA	AGG	GCC	GCA	AAA	GGA	GAT	TTG	2160
Leu	Ile	Leu	Thr	Thr	Gly	Asp	Ile	Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu	
705					710					715					720	
GTA	CCA	TTT	GCA	CTC	TTA	CAC	TTA	TTG	CAT	TGT	TTG	TTA	TCC	AAG	TCA	2208
Val	Pro	Phe	Ala	Leu	Leu	His	Leu	Leu	His	Cys	Leu	Leu	Ser	Lys	Ser	
				725					730					735		
GCA	TCT	GTC	TCT	GGA	GCA	GCA	TAC	ACA	GAA	ATT	AGA	GCT	CTG	GTT	GCA	2256
Ala	Ser	Val	Ser	Gly	Ala	Ala	Tyr	Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala	
			740					745					750			
GCT	AAA	AGT	GTT	AAA	CTG	CAA	AGT	TTT	TTC	AGC	CAG	TAT	AAG	AAA	CCC	2304
Ala	Lys	Ser	Val	Lys	Leu	Gln	Ser	Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro	
		755					760					765				
ATC	TGT	CAG	TTT	TTG	GTA	GAA	TCC	CTT	CAC	TCT	AGT	CAG	ATG	ACA	GCA	2352
Ile	Cys	Gln	Phe	Leu	Val	Glu	Ser	Leu	His	Ser	Ser	Gln	Met	Thr	Ala	
	770					775					780					
CTT	CCG	AAT	ACT	CCA	TGC	CAG	AAT	GCT	GAC	GTG	CGA	AAA	CAA	GAT	GTG	2400
Leu	Pro	Asn	Thr	Pro	Cys	Gln	Asn	Ala	Asp	Val	Arg	Lys	Gln	Asp	Val	
785					790					795					800	
GCT	CAC	CAG	AGA	GAA	ATG	GCT	TTA	AAT	ACG	TTG	TCT	GAA	ATT	GCC	AAC	2448
Ala	His	Gln	Arg	Glu	Met	Ala	Leu	Asn	Thr	Leu	Ser	Glu	Ile	Ala	Asn	
				805					810					815		
GTT	TTC	GAC	TTT	CCT	GAT	CTT	AAT	CGT	TTT	CTT	ACT	AGG	ACA	TTA	CAA	2496
Val	Phe	Asp	Phe	Pro	Asp	Leu	Asn	Arg	Phe	Leu	Thr	Arg	Thr	Leu	Gln	
			820					825					830			
GTT	CTA	CTA	CCT	GAT	CTT	GCT	GCC	AAA	GCA	AGC	CCT	GCA	GCT	TCT	GCT	2544
Val	Leu	Leu	Pro	Asp	Leu	Ala	Ala	Lys	Ala	Ser	Pro	Ala	Ala	Ser	Ala	
			835				840					845				
CTC	ATT	CGA	ACT	TTA	GGA	AAA	CAA	TTA	AAT	GTC	AAT	CGT	AGA	GAG	ATT	2592
Leu	Ile	Arg	Thr	Leu	Gly	Lys	Gln	Leu	Asn	Val	Asn	Arg	Arg	Glu	Ile	
	850					855					860					
TTA	ATA	AAC	AAC	TTC	AAA	TAT	ATT	TTT	TCT	CAT	TTG	GTC	TGT	TCT	TGT	2640
Leu	Ile	Asn	Asn	Phe	Lys	Tyr	Ile	Phe	Ser	His	Leu	Val	Cys	Ser	Cys	
865					870					875					880	
TCC	AAA	GAT	GAA	TTA	GAA	CGT	GCC	CTT	CAT	TAT	CTG	AAG	AAT	GAA	ACA	2688
Ser	Lys	Asp	Glu	Leu	Glu	Arg	Ala	Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr	
				885					890					895		
GAA	ATT	GAA	CTG	GGG	AGC	CTG	TTG	AGA	CAA	GAT	TTC	CAA	GGA	TTG	CAT	2736
Glu	Ile	Glu	Leu	Gly	Ser	Leu	Leu	Arg	Gln	Asp	Phe	Gln	Gly	Leu	His	
			900					905					910			
AAT	GAA	TTA	TTG	CTG	CGT	ATT	GGA	GAA	CAC	TAT	CAA	CAG	GTT	TTT	AAT	2784
Asn	Glu	Leu	Leu	Leu	Arg	Ile	Gly	Glu	His	Tyr	Gln	Gln	Val	Phe	Asn	
		915					920					925				
GGT	TTG	TCA	ATA	CTT	GCC	TCA	TTT	GCA	TCC	AGT	GAT	GAT	CCA	TAT	CAG	2832
Gly	Leu	Ser	Ile	Leu	Ala	Ser	Phe	Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln	
	930					935					940					
GGC	CCG	AGA	GAT	ATC	ATA	TCA	CCT	GAA	CTG	ATG	GCT	GAT	TAT	TTA	CAA	2880
Gly	Pro	Arg	Asp	Ile	Ile	Ser	Pro	Glu	Leu	Met	Ala	Asp	Tyr	Leu	Gln	
945					950					955					960	
CCC	AAA	TTG	TTG	GGC	ATT	TTG	GCT	TTT	TTT	AAC	ATG	CAG	TTA	CTG	AGC	2928
Pro	Lys	Leu	Leu	Gly	Ile	Leu	Ala	Phe	Phe	Asn	Met	Gln	Leu	Leu	Ser	
				965					970					975		

-117-

TCT AGT GTT GGC ATT GAA GAT AAG AAA ATG GCC TTG AAC AGT TTG ATG Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met 980 985 990	2976
TCT TTG ATG AAG TTA ATG GGA CCC AAA CAT GTC AGT TCT GTG AGG GTG Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val 995 1000 1005	3024
AAG ATG ATG ACC ACA CTG AGA ACT GGC CTT CGA TTC AAG GAT GAT TTT Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe 1010 1015 1020	3072
CCT GAA TTG TGT TGC AGA GCT TGG GAC TGC TTT GTT CGC TGC CTG GAT Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp 1025 1030 1035 1040	3120
CAT GCT TGT CTG GGC TCC CTT CTC AGT CAT GTA ATA GTA GCT TTG TTA His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu 1045 1050 1055	3168
CCT CTT ATA CAC ATC CAG CCT AAA GAA ACT GCA GCT ATC TTC CAC TAC Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr 1060 1065 1070	3216
CTC ATA ATT GAA AAC AGG GAT GCT GTG CAA GAT TTT CTT CAT GAA ATA Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile 1075 1080 1085	3264
TAT TTT TTA CCT GAT CAT CCA GAA TTA AAA AAG ATA AAA GCC GTT CTC Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu 1090 1095 1100	3312
CAG GAA TAC AGA AAG GAG ACC TCT GAG AGC ACT GAT CTT CAG ACA ACT Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr 1105 1110 1115 1120	3360
CTT CAG CTC TCT ATG AAG GCC ATT CAA CAT GAA AAT GTC GAT GTT CGT Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg 1125 1130 1135	3408
ATT CAT GCT CTT ACA AGC TTG AAG GAA ACC TTG TAT AAA AAT CAG GAA Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu 1140 1145 1150	3456
AAA CTG ATA AAG TAT GCA ACA GAC AGT GAA ACA GTA GAA CCT ATT ATC Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile 1155 1160 1165	3504
TCA CAG TTG GTG ACA GTG CTT TTG AAA GGT TGC CAA GAT GCA AAC TCT Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser 1170 1175 1180	3552
CAA GCT CGG TTG CTC TGT GGG GAA TGT TTA GGG GAA TTG GGG GCG ATA Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile 1185 1190 1195 1200	3600
GAT CCA GGT CGA TTA GAT TTC TCA ACA ACT GAA ACT CAA GGA AAA GAT Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp 1205 1210 1215	3648
TTT ACA TTT GTG ACT GGA GTA GAA GAT TCA AGC TTT GCC TAT GGA TTA Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu 1220 1225 1230	3696
TTG ATG GAG CTA ACA AGA GCT TAC CTT GCG TAT GCT GAT AAT AGC CGA Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg 1235 1240 1245	3744

-118-

GCT CAA GAT TCA GCT GCC TAT GCC ATT CAG GAG TTG CTT TCT ATT TAT Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr 1250 1255 1260	3792
GAC TGT AGA GAG ATG GAG ACC AAC GGC CCA GGT CAC CAA TTG TGG AGG Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg 1265 1270 1275 1280	3840
AGA TTT CCT GAG CAT GTT CGG GAA ATA CTA GAA CCT CAT CTA AAT ACC Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr 1285 1290 1295	3888
AGA TAC AAG AGT TCT CAG AAG TCA ACC GAT TGG TCT GGA GTA AAG AAG Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys 1300 1305 1310	3936
CCA ATT TAC TTA AGT AAA TTG GGT AGT AAC TTT GCA GAA TGG TCA GCA Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala 1315 1320 1325	3984
TCT TGG GCA GGT TAT CTT ATT ACA AAG GTT CGA CAT GAT CTT GCC AGT Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser 1330 1335 1340	4032
AAA ATT TTC ACC TGC TGT AGC ATT ATG ATG AAG CAT GAT TTC AAA GTG Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val 1345 1350 1355 1360	4080
ACC ATC TAT CTT CTT CCA CAT ATT CTG GTG TAT GTC TTA CTG GGT TGT Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys 1365 1370 1375	4128
AAT CAA GAA GAT CAG CAG GAG GTT TAT GCA GAA ATT ATG GCA GTT CTA Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu 1380 1385 1390	4176
AAG CAT GAC GAT CAG CAT ACC ATA AAT ACC CAA GAC ATT GCA TCT GAT Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp 1395 1400 1405	4224
CTG TGT CAA CTC AGT ACA CAG ACT GTG TTC TCC ATG CTT GAC CAT CTC Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu 1410 1415 1420	4272
ACA CAG TGG GCA AGG CAC AAA TTT CAG GCA CTG AAA GCT GAG AAA TGT Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys 1425 1430 1435 1440	4320
CCA CAC AGC AAA TCA AAC AGA AAT AAG GTA GAC TCA ATG GTA TCT ACT Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr 1445 1450 1455	4368
GTG GAT TAT GAA GAC TAT CAG AGT GTA ACC CGT TTT CTA GAC CTC ATA Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile 1460 1465 1470	4416
CCC CAG GAT ACT CTG GCA GTA GCT TCC TTT CGC TCC AAA GCA TAC ACA Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr 1475 1480 1485	4464
CGA GCT GTA ATG CAC TTT GAA TCA TTT ATT ACA GAA AAG AAG CAA AAT Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn 1490 1495 1500	4512
ATT CAG GAA CAT CTT GGA TTT TTA CAG AAA TTG TAT GCT GCT ATG CAT Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His 1505 1510 1515 1520	4560

-119-

GAA CCT GAT GGA GTG GCC GGA GTC AGT GCA ATT AGA AAG GCA GAA CCA Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg Lys Ala Glu Pro 1525 1530 1535	4608
TCT CTA AAA GAA CAG ATC CTT GAA CAT GAA AGC CTT GGC TTG CTG AGG Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg 1540 1545 1550	4656
GAT GCC ACT GCT TGT TAT GAC AGG GCT ATT CAG CTA GAA CCA GAC CAG Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln 1555 1560 1565	4704
ATC ATT CAT TAC CAT GGT GTA GTA AAG TCC ATG TTA GGT CTT GGT CAG Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln 1570 1575 1580	4752
CTG TCT ACT GTT ATC ACT CAG GTG AAT GGA GTG CAT GCT AAC AGG TCC Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser 1585 1590 1595 1600	4800
GAG TGG ACA GAT GAA TTA AAC ACG TAC AGA GTG GAA GCA GCT TGG AAA Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys 1605 1610 1615	4848
TTG TCA CAG TGG GAT TTG GTG GAA AAC TAT TTG GCA GCA GAT GGA AAA Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys 1620 1625 1630	4896
TCT ACA ACA TGG AGT GTC AGA CTG GGA CAG CTA TTA TTA TCA GCC AAA Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys 1635 1640 1645	4944
AAA AGA GAT ATC ACA GCT TTT TAT GAC TCA CTG AAA CTA GTG AGA GCA Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala 1650 1655 1660	4992
GAA CAA ATT GTA CCT CTT TCA GCT GCA AGC TTT GAA AGA GGC TCC TAC Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr 1665 1670 1675 1680	5040
CAA CGA GGA TAT GAA TAT ATT GTG AGA TTG CAC ATG TTA TGT GAG TTG Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu 1685 1690 1695	5088
GAG CAT AGC ATC AAA CCA CTT TTC CAG CAT TCT CCA GGT GAC AGT TCT Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser 1700 1705 1710	5136
CAA GAA GAT TCT CTA AAC TGG GTA GCT CGA CTA GAA ATG ACC CAG AAT Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn 1715 1720 1725	5184
TCC TAC AGA GCC AAG GAG CCT ATC CTG GCT CTC CGG AGG GCT TTA CTA Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu 1730 1735 1740	5232
AGC CTC AAC AAA AGA CCA GAT TAC AAT GAA ATG GTT GGA GAA TGC TGG Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp 1745 1750 1755 1760	5280
CTG CAG AGT GCC AGG GTA GCT AGA AAG GCT GGT CAC CAC CAG ACA GCC Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala 1765 1770 1775	5328
TAC AAT GCT CTC CTT AAT GCA GGG GAA TCA CGA CTC GCT GAA CTG TAC Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr 1780 1785 1790	5376

-120-

GTG GAA AGG GCA AAG TGG CTC TGG TCC AAG GGT GAT GTT CAC CAG GCA Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala 1795 1800 1805	5424
CTA ATT GTT CTT CAA AAA GGT GTT GAA TTA TGT TTT CCT GAA AAT GAA Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu 1810 1815 1820	5472
ACC CCA CCT GAG GGT AAG AAC ATG TTA ATC CAT GGT CGA GCT ATG CTA Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu 1825 1830 1835 1840	5520
CTA GTG GGC CGA TTT ATG GAA GAA ACA GCT AAC TTT GAA AGC AAT GCA Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala 1845 1850 1855	5568
ATT ATG AAA AAA TAT AAG GAT GTG ACC GCG TGC CTG CCA GAA TGG GAG Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu 1860 1865 1870	5616
GAT GGG CAT TTT TAC CTT GCC AAG TAC TAT GAC AAA TTG ATG CCC ATG Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met 1875 1880 1885	5664
GTC ACA GAC AAC AAA ATG GAA AAG CAA GGT GAT CTC ATC CGG TAT ATA Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile 1890 1895 1900	5712
GTT CTT CAT TTT GGC AGA TCT CTA CAA TAT GGA AAT CAG TTC ATA TAT Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr 1905 1910 1915 1920	5760
CAG TCA ATG CCA CGA ATG TTA ACT CTA TGG CTT GAT TAT GGT ACA AAG Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys 1925 1930 1935	5808
GCA TAT GAA TGG GAA AAA GCT GGC CGC TCC GAT CGT GTA CAA ATG AGG Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg 1940 1945 1950	5856
AAT GAT TTG GGT AAA ATA AAC AAG GTT ATC ACA GAG CAT ACA AAC TAT Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr 1955 1960 1965	5904
TTA GCT CCA TAT CAA TTT TTG ACT GCT TTT TCA CAA TTG ATC TCT CGA Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg 1970 1975 1980	5952
ATT TGT CAT TCT CAC GAT GAA GTT TTT GTT GTG CTT GAT GGA AAT AAT Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn 1985 1990 1995 2000	6000
AGC CAA GTA TTT CTA GCC TAT CCT CAA CAA GCA ATG TGG ATG ATG ACA Ser Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr 2005 2010 2015	6048
GCT GTG TCA AAG TCA TCT TAT CCC ATG CGT GTG AAC AGA TGC AAG GAA Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu 2020 2025 2030	6096
ATC CTC AAT AAA GCT ATT CAT ATG AAA AAA TCC TTA GAG AAG TTT GTT Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val 2035 2040 2045	6144
GGA GAT GCA ACT CGC CTA ACA GAT AAG CTT CTA GAA TTG TGC AAT AAA Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys 2050 2055 2060	6192

-121-

CCG GTT GAT GGA AGT AGT TCC ACA TTA AGC ATG AGC ACT CAT TTT AAA Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys 2065 2070 2075 2080	6240
ATG CTT AAA AAG CTG GTA GAA GAA GCA ACA TTT AGT GAA ATC CTC ATT Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile 2085 2090 2095	6288
CCT CTA CAA TCA GTC ATG ATA CCT ACA CTT CCA TCA ATT CTG GGT ACC Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr 2100 2105 2110	6336
CAT GCT AAC CAT GCT AGC CAT GAA CCA TTT CCT GGA CAT TGG GCC TAT His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr 2115 2120 2125	6384
ATT GCA GGG TTT GAT GAT ATG GTG GAA ATT CTT GCT TCT CTT CAG AAA Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys 2130 2135 2140	6432
CCA AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG TTC TAC ATC ATG Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met 2145 2150 2155 2160	6480
ATG TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT AGA CTA ATG GAA Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu 2165 2170 2175	6528
TTC AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT GCA GAG TCT CGT Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg 2180 2185 2190	6576
AGA AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT CCA CTA AAT GAT Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp 2195 2200 2205	6624
GAA TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT GGT TTG AGA CCT Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro 2210 2215 2220	6672
ATT CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT ATG ACA GGA AAA Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys 2225 2230 2235 2240	6720
GAA CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT TTA TCT GAA AAA Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys 2245 2250 2255	6768
CTC AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT CCT CCT ATT TTT Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Ile Phe 2260 2265 2270	6816
CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA TCA TGG TAC AGT His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser 2275 2280 2285	6864
AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA ATG GTT GGT Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly 2290 2295 2300	6912
TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT CTC TTT GAT Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp 2305 2310 2315 2320	6960
TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT CTT TTC AAT Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn 2325 2330 2335	7008

-122-

AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT CGC CTG ACT Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr 2340 2345 2350	7056
CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG GGT CTT TTT His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe 2355 2360 2365	7104
CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT CAG CGA GAG Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu 2370 2375 2380	7152
CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT CTT GTG GAA Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu 2385 2390 2395 2400	7200
TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG AAT GAA ACT Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr 2405 2410 2415	7248
GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT GAC ATT GAG Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu 2420 2425 2430	7296
CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG ACA GGA CTG Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu 2435 2440 2445	7344
CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA GAA GCT ACT Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr 2450 2455 2460	7392
GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT CCA TAT ATG Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met 2465 2470 2475 2480	7440
TGAAATGAAA TTATGTAAAA GAATATGTTA ATAATCTAAA AGTAAAAAAA AAAAAAAAAA	7500
AA	7502

## (2) INFORMATION FOR SEQ ID NO:26:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2480 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser 1 5 10 15
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu 20 25 30
Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr Leu Leu Met Val 35 40 45
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu 50 55 60
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile 65 70 75 80



Lys	Ser	Leu	Ala	Ile	Ser	Phe	Leu	Thr	Glu	Leu	Phe	Gln	Leu	Gly	Gly	
				85					90					95		
Leu	Pro	Ala	Gln	Pro	Ala	Ser	Thr	Phe	Phe	Ser	Ser	Phe	Leu	Glu	Leu	
				100					105					110		
Leu	Lys	His	Leu	Val	Glu	Met	Asp	Thr	Asp	Gln	Leu	Lys	Leu	Tyr	Glu	
				115					120					125		
Glu	Pro	Leu	Ser	Lys	Leu	Ile	Lys	Thr	Leu	Phe	Pro	Phe	Glu	Ala	Glu	
				130					135					140		
Ala	Tyr	Arg	Asn	Ile	Glu	Pro	Val	Tyr	Leu	Asn	Met	Leu	Leu	Glu	Lys	
				145					150					155		
Leu	Cys	Val	Met	Phe	Glu	Asp	Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	
				165					170					175		
Leu	Leu	Lys	Ala	Ala	Leu	Cys	His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	
				180					185					190		
Val	Pro	Ala	Gly	Tyr	Glu	Ser	Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	
				195					200					205		
Arg	Asn	Ile	Cys	Lys	Ala	Leu	Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	
				210					215					220		
Ala	Glu	Tyr	Leu	Leu	Gly	Pro	Leu	Tyr	Ala	Ala	Leu	Lys	Met	Glu	Ser	
				225					230					235		
Met	Glu	Ile	Ile	Glu	Glu	Ile	Gln	Cys	Gln	Thr	Gln	Gln	Glu	Asn	Leu	
				245					250					255		
Ser	Ser	Asn	Ser	Asp	Gly	Ile	Ser	Pro	Lys	Arg	Arg	Arg	Leu	Ser	Ser	
				260					265					270		
Ser	Leu	Asn	Pro	Ser	Lys	Arg	Ala	Pro	Lys	Gln	Thr	Glu	Glu	Ile	Lys	
				275					280					285		
His	Val	Asp	Met	Asn	Gln	Lys	Ser	Ile	Leu	Trp	Ser	Ala	Leu	Lys	Gln	
				290					295					300		
Lys	Ala	Glu	Ser	Leu	Gln	Ile	Ser	Leu	Glu	Tyr	Ser	Gly	Leu	Lys	Asn	
				305					310					315		
Pro	Val	Ile	Glu	Met	Leu	Glu	Gly	Ile	Ala	Val	Val	Leu	Gln	Leu	Thr	
				325					330					335		
Ala	Leu	Cys	Thr	Val	His	Cys	Ser	His	Gln	Asn	Met	Asn	Cys	Arg	Thr	
				340					345					350		
Phe	Lys	Asp	Cys	Gln	His	Lys	Ser	Lys	Lys	Lys	Pro	Ser	Val	Val	Ile	
				355					360					365		
Thr	Trp	Met	Ser	Leu	Asp	Phe	Tyr	Thr	Lys	Val	Leu	Lys	Ser	Cys	Arg	
				370					375					380		
Ser	Leu	Leu	Glu	Ser	Val	Gln	Lys	Leu	Asp	Leu	Glu	Ala	Thr	Ile	Asp	
				385					390					395		
Lys	Val	Val	Lys	Ile	Tyr	Asp	Ala	Leu	Ile	Tyr	Met	Gln	Val	Asn	Ser	
				405					410					415		
Ser	Phe	Glu	Asp	His	Ile	Leu	Glu	Asp	Leu	Cys	Gly	Met	Leu	Ser	Leu	
				420					425					430		

Pro	Trp	Ile	Tyr	Ser	His	Ser	Asp	Asp	Gly	Cys	Leu	Lys	Leu	Thr	Thr	
		435					440					445				
Phe	Ala	Ala	Asn	Leu	Leu	Thr	Leu	Ser	Cys	Arg	Ile	Ser	Asp	Ser	Tyr	
	450					455					460					
Ser	Pro	Gln	Ala	Gln	Ser	Arg	Cys	Val	Phe	Leu	Leu	Thr	Leu	Phe	Pro	
465					470					475					480	
Arg	Arg	Ile	Phe	Leu	Glu	Trp	Arg	Thr	Ala	Val	Tyr	Asn	Trp	Ala	Leu	
				485					490					495		
Gln	Ser	Ser	His	Glu	Val	Ile	Arg	Ala	Ser	Cys	Val	Ser	Gly	Phe	Phe	
			500					505					510			
Ile	Leu	Leu	Gln	Gln	Gln	Asn	Ser	Cys	Asn	Arg	Val	Pro	Lys	Ile	Leu	
		515					520					525				
Ile	Asp	Lys	Val	Lys	Asp	Asp	Ser	Asp	Ile	Val	Lys	Lys	Glu	Phe	Ala	
	530					535					540					
Ser	Ile	Leu	Gly	Gln	Leu	Val	Cys	Thr	Leu	His	Gly	Met	Phe	Tyr	Leu	
545					550					555					560	
Thr	Ser	Ser	Leu	Thr	Glu	Pro	Phe	Ser	Glu	His	Gly	His	Val	Asp	Leu	
				565					570					575		
Phe	Cys	Arg	Asn	Leu	Lys	Ala	Thr	Ser	Gln	His	Glu	Cys	Ser	Ser	Ser	
			580					585					590			
Gln	Leu	Lys	Ala	Ser	Val	Cys	Lys	Pro	Phe	Leu	Phe	Leu	Leu	Lys	Lys	
		595					600					605				
Lys	Ile	Pro	Ser	Pro	Val	Lys	Leu	Ala	Phe	Ile	Asp	Asn	Leu	His	His	
	610					615					620					
Leu	Cys	Lys	His	Leu	Asp	Phe	Arg	Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala	
625					630					635					640	
Val	Leu	Gly	Thr	Leu	Leu	Asn	Leu	Met	Glu	Asp	Pro	Asp	Lys	Asp	Val	
				645					650					655		
Arg	Val	Ala	Phe	Ser	Gly	Asn	Ile	Lys	His	Ile	Leu	Glu	Ser	Leu	Asp	
			660					665					670			
Ser	Glu	Asp	Gly	Phe	Ile	Lys	Glu	Leu	Phe	Val	Leu	Arg	Met	Lys	Glu	
		675					680					685				
Ala	Tyr	Thr	His	Ala	Gln	Ile	Ser	Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr	
	690					695					700					
Leu	Ile	Leu	Thr	Thr	Gly	Asp	Ile	Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu	
705					710					715					720	
Val	Pro	Phe	Ala	Leu	Leu	His	Leu	Leu	His	Cys	Leu	Leu	Ser	Lys	Ser	
				725					730					735		
Ala	Ser	Val	Ser	Gly	Ala	Ala	Tyr	Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala	
			740					745					750			
Ala	Lys	Ser	Val	Lys	Leu	Gln	Ser	Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro	
		755					760					765				
Ile	Cys	Gln	Phe													

-125-

Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val  
 785 790 795 800  
 Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn  
 805 810 815  
 Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln  
 820 825 830  
 Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala  
 835 840 845  
 Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile  
 850 855 860  
 Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys  
 865 870 875 880  
 Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu Lys Asn Glu Thr  
 885 890 895  
 Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe Gln Gly Leu His  
 900 905 910  
 Asn Glu Leu Leu Leu Arg Ile Gly Glu His Tyr Gln Gln Val Phe Asn  
 915 920 925  
 Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp Asp Pro Tyr Gln  
 930 935 940  
 Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln  
 945 950 955 960  
 Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser  
 965 970 975  
 Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met  
 980 985 990  
 Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val  
 995 1000 1005  
 Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe  
 1010 1015 1020  
 Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp  
 1025 1030 1035 1040  
 His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu  
 1045 1050 1055  
 Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr  
 1060 1065 1070  
 Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile  
 1075 1080 1085  
 Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu  
 1090 1095 1100  
 Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr  
 1105 1110 1115 1120  
 Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg  
 1125 1130 1135

-126-

Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu  
 1140 1145 1150  
 Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile  
 1155 1160 1165  
 Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser  
 1170 1175 1180  
 Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile  
 1185 1190 1195 1200  
 Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp  
 1205 1210 1215  
 Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu  
 1220 1225 1230  
 Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg  
 1235 1240 1245  
 Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr  
 1250 1255 1260  
 Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg  
 1265 1270 1275 1280  
 Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr  
 1285 1290 1295  
 Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys  
 1300 1305 1310  
 Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala  
 1315 1320 1325  
 Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser  
 1330 1335 1340  
 Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val  
 1345 1350 1355 1360  
 Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys  
 1365 1370 1375  
 Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu  
 1380 1385 1390  
 Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp  
 1395 1400 1405  
 Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu  
 1410 1415 1420  
 Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys  
 1425 1430 1435 1440  
 Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr  
 1445 1450 1455  
 Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile  
 1460 1465 1470  
 Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr  
 1475 1480 1485

-127-

Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn  
 1490 1495 1500  
 Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His  
 1505 1510 1515 1520  
 Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg Lys Ala Glu Pro  
 1525 1530 1535  
 Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg  
 1540 1545 1550  
 Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln  
 1555 1560 1565  
 Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln  
 1570 1575 1580  
 Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser  
 1585 1590 1595 1600  
 Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys  
 1605 1610 1615  
 Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys  
 1620 1625 1630  
 Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys  
 1635 1640 1645  
 Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala  
 1650 1655 1660  
 Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr  
 1665 1670 1675 1680  
 Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu  
 1685 1690 1695  
 Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser  
 1700 1705 1710  
 Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn  
 1715 1720 1725  
 Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu  
 1730 1735 1740  
 Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp  
 1745 1750 1755 1760  
 Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala  
 1765 1770 1775  
 Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr  
 1780 1785 1790  
 Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala  
 1795 1800 1805  
 Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu  
 1810 1815 1820  
 Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu  
 1825 1830 1835 1840

-128-

Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala  
 1845 1850 1855  
 Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu  
 1860 1865 1870  
 Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met  
 1875 1880 1885  
 Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile  
 1890 1895 1900  
 Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr  
 1905 1910 1915 1920  
 Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys  
 1925 1930 1935  
 Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg  
 1940 1945 1950  
 Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr  
 1955 1960 1965  
 Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg  
 1970 1975 1980  
 Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Asp Gly Asn Asn  
 1985 1990 1995 2000  
 Ser Gln Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr  
 2005 2010 2015  
 Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu  
 2020 2025 2030  
 Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val  
 2035 2040 2045  
 Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys  
 2050 2055 2060  
 Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys  
 2065 2070 2075 2080  
 Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile  
 2085 2090 2095  
 Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr  
 2100 2105 2110  
 His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr  
 2115 2120 2125  
 Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys  
 2130 2135 2140  
 Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met  
 2145 2150 2155 2160  
 Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu  
 2165 2170 2175  
 Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg  
 2180 2185 2190

-129-

Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp  
 2195 2200 2205  
 Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro  
 2210 2215 2220  
 Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys  
 2225 2230 2235 2240  
 Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys  
 2245 2250 2255  
 Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro Ile Phe  
 2260 2265 2270  
 His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser  
 2275 2280 2285  
 Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly  
 2290 2295 2300  
 Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp  
 2305 2310 2315 2320  
 Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn  
 2325 2330 2335  
 Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr  
 2340 2345 2350  
 His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe  
 2355 2360 2365  
 Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu  
 2370 2375 2380  
 Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu  
 2385 2390 2395 2400  
 Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr  
 2405 2410 2415  
 Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu  
 2420 2425 2430  
 Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu  
 2435 2440 2445  
 Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr  
 2450 2455 2460  
 Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2465 2470 2475 2480

## (2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 878 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

-130-

ATCCATTGTG TTGGAAAGGA ATGATGAATG TGGGATTATT GAATGGGTGA ACAATACTGC	60
TGGCTTGAGA CCTATTCTGA CCAAAATATA TAAAGAAAAG GGAGTGTATA TGACAGGAAA	120
GGAGCTTCGC CAGTGTATGC TACCAAAGTC AGCAGCTTTA TCTGAAAAAC TCAAAGTATT	180
CCAAGAATTA CTCCTGCCCA GGCATCCTCC TGTTTTTTCAT GAGTGGTTTC TGAGAACATT	240
CCCTGATCCT ACATCATGGT ACAGTAGCAG ATCTGCATAT TGCCGCTCTA CTGCAGTCAT	300
GTCAATGGTT GGCTACATCC TGGGGCTTGG AGACCGTCAT GGTGAAAACA TTCTTTTTGA	360
CTCTTTTCACT GGTGAATGTG TACATGTAGA TTTCAACTGT CTTTTTAATA AGGGAGAAAC	420
GTTTGAAGTT CCGGAAATTG TACCATTTTCG ACTGACTCAT AATATGGTTA ATGGAATGGG	480
TCCTATGGGA ACAGAGGGTC TATTTTGAAG AGCATGTGAA GTTACACTGA GACTGATGAG	540
GGATCAGAGA GAACCTTTAA TGAGTGTCTT AAAGACTTTT CTACACGATC CTCTAGTGGA	600
GTGGAGTAAA CCAGTGAAAG GACACTCCAA AGCACCCTG AATGAAACCG GGGAAGTTGT	660
CAATGAGAAG GCCAAGACCC ATGTTCTTGA CATTGAACAA CGACTACAAG GTGTGATCAA	720
AACCCGAAAT AGAGTAACAG GGCTGCCATT ATCTATTGAA GGACATGTGC ATTACCTCAT	780
ACAAGAAGCT ACTGATGAAA ACTTACTCTG TCAGATGTAC CTTGGTTGGA CCCCATATAT	840
GTAAAATAAA ATTATTTCAA AGAAAAAAAA AAAAAAAAAA	878

## (2) INFORMATION FOR SEQ ID NO:28:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7935 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..7932

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

ATG GGG GAA CAT GGC CTG GAG CTG GCT TCC ATG ATC CCC GCC CTG CGG	48
Met Gly Glu His Gly Leu Glu Leu Ala Ser Met Ile Pro Ala Leu Arg	
1 5 10 15	
GAG CTG GGC AGT GCC ACA CCA GAG GAA TAT AAT ACA GTT GTA CAG AAG	96
Glu Leu Gly Ser Ala Thr Pro Glu Tyr Asn Thr Val Val Gln Lys	
20 25 30	



-131-

CCA	AGA	CAA	ATT	CTG	TGT	CAA	TTC	ATT	GAC	CGG	ATA	CTT	ACA	GAT	GTA	144
Pro	Arg	Gln	Ile	Leu	Cys	Gln	Phe	Ile	Asp	Arg	Ile	Leu	Thr	Asp	Val	
		35					40					45				
AAT	GTT	GTT	GCT	GTA	GAA	CTT	GTA	AAG	AAA	ACT	GAC	TCT	CAG	CCA	ACC	192
Asn	Val	Val	Ala	Val	Glu	Leu	Val	Lys	Lys	Thr	Asp	Ser	Gln	Pro	Thr	
	50					55					60					
TCC	GTG	ATG	TTG	CTT	GAT	TTC	ATC	CAG	CAT	ATC	ATG	AAA	TCC	TCC	CCA	240
Ser	Val	Met	Leu	Leu	Asp	Phe	Ile	Gln	His	Ile	Met	Lys	Ser	Ser	Pro	
	65				70					75					80	
CTT	ATG	TTT	GTA	AAT	GTG	AGT	GGA	AGC	CAT	GAG	CGC	AAA	GGC	AGT	TGT	288
Leu	Met	Phe	Val	Asn	Val	Ser	Gly	Ser	His	Glu	Arg	Lys	Gly	Ser	Cys	
				85					90					95		
ATT	GAA	TTC	AGT	AAT	TGG	ATC	ATA	ACG	AGA	CTT	CTG	CGG	ATT	GCA	GCA	336
Ile	Glu	Phe	Ser	Asn	Trp	Ile	Ile	Thr	Arg	Leu	Leu	Arg	Ile	Ala	Ala	
			100					105					110			
ACT	CCC	TCC	TGT	CAT	TTG	TTA	CAC	AAG	AAA	ATC	TGT	GAA	GTC	ATC	TGT	384
Thr	Pro	Ser	Cys	His	Leu	Leu	His	Lys	Lys	Ile	Cys	Glu	Val	Ile	Cys	
		115					120					125				
TCA	TTA	TTA	TTT	CTT	TTT	AAA	AGC	AAG	AGT	CCT	GCT	ATT	TTT	GGG	GTA	432
Ser	Leu	Leu	Phe	Leu	Phe	Lys	Ser	Lys	Ser	Pro	Ala	Ile	Phe	Gly	Val	
	130					135					140					
CTC	ACA	AAA	GAA	TTA	TTA	CAA	CTT	TTT	GAA	GAC	TTG	GTT	TAC	CTC	CAT	480
Leu	Thr	Lys	Glu	Leu	Leu	Gln	Leu	Phe	Glu	Asp	Leu	Val	Tyr	Leu	His	
	145					150				155					160	
AGA	AGA	AAT	GTG	ATG	GGT	CAT	GCT	GTG	GAA	TGG	CCA	GTG	GTC	ATG	AGC	528
Arg	Arg	Asn	Val	Met	Gly	His	Ala	Val	Glu	Trp	Pro	Val	Val	Met	Ser	
				165					170					175		
CGA	TTT	TTA	AGT	CAA	TTA	GAT	GAA	CAC	ATG	GGA	TAT	TTA	CAA	TCA	GCT	576
Arg	Phe	Leu	Ser	Gln	Leu	Asp	Glu	His	Met	Gly	Tyr	Leu	Gln	Ser	Ala	
			180					185					190			
CCT	TTG	CAG	TTG	ATG	AGT	ATG	CAA	AAT	TTA	GAA	TTT	ATT	GAA	GTC	ACT	624
Pro	Leu	Gln	Leu	Met	Ser	Met	Gln	Asn	Leu	Glu	Phe	Ile	Glu	Val	Thr	
		195					200					205				
TTA	TTA	ATG	GTT	CTT	ACT	CGT	ATT	ATT	GCA	ATT	GTG	TTT	TTT	AGA	AGG	672
Leu	Leu	Met	Val	Leu	Thr	Arg	Ile	Ile	Ala	Ile	Val	Phe	Phe	Arg	Arg	
	210					215					220					
CAA	GAA	CTC	TTA	CTT	TGG	CAG	ATA	GGT	TGT	GTT	CTG	CTA	GAG	TAT	GGT	720
Gln	Glu	Leu	Leu	Leu	Trp	Gln	Ile	Gly	Cys	Val	Leu	Leu	Glu	Tyr	Gly	
	225				230				235						240	
AGT	CCA	AAA	ATT	AAA	TCC	CTA	GCA	ATT	AGC	TTT	TTA	ACA	GAA	CTT	TTT	768
Ser	Pro	Lys	Ile	Lys	Ser	Leu	Ala	Ile	Ser	Phe	Leu	Thr	Glu	Leu	Phe	
				245					250					255		
CAG	CTT	GGA	GGA	CTA	CCA	GCA	CAA	CCA	GCT	AGC	ACT	TTT	TTC	AGC	TCA	816
Gln	Leu	Gly	Gly	Leu	Pro	Ala	Gln	Pro	Ala	Ser	Thr	Phe	Phe	Ser	Ser	
			260					265					270			
TTT	TTG	GAA	TTA	TTA	AAA	CAC	CTT	GTA	GAA	ATG	GAT	ACT	GAC	CAA	TTG	864
Phe	Leu	Glu	Leu	Leu	Lys	His	Leu	Val	Glu	Met	Asp	Thr	Asp	Gln	Leu	
		275					280					285				
AAA	CTC	TAT	GAA	GAG	CCA	TTA	TCA	AAG	CTG	ATA	AAG	ACA	CTA	TTT	CCC	912
Lys	Leu	Tyr	Glu	Glu	Pro	Leu	Ser	Lys	Leu	Ile	Lys	Thr	Leu	Phe	Pro	
	290					295					300					

-132-

TTT	GAA	GCA	GAA	GCT	TAT	AGA	AAT	ATT	GAA	CCT	GTC	TAT	TTA	AAT	ATG	960
Phe	Glu	Ala	Glu	Ala	Tyr	Arg	Asn	Ile	Glu	Pro	Val	Tyr	Leu	Asn	Met	
305					310					315					320	
CTG	CTG	GAA	AAA	CTC	TGT	GTC	ATG	TTT	GAA	GAC	GGT	GTG	CTC	ATG	CGG	1008
Leu	Leu	Glu	Lys	Leu	Cys	Val	Met	Phe	Glu	Asp	Gly	Val	Leu	Met	Arg	
				325					330					335		
CTT	AAG	TCT	GAT	TTG	CTA	AAA	GCA	GCT	TTG	TGC	CAT	TTA	CTG	CAG	TAT	1056
Leu	Lys	Ser	Asp	Leu	Leu	Lys	Ala	Ala	Leu	Cys	His	Leu	Leu	Gln	Tyr	
			340					345					350			
TTC	CTT	AAA	TTT	GTG	CCA	GCT	GGG	TAT	GAA	TCT	GCT	TTA	CAA	GTC	AGG	1104
Phe	Leu	Lys	Phe	Val	Pro	Ala	Gly	Tyr	Glu	Ser	Ala	Leu	Gln	Val	Arg	
		355					360					365				
AAG	GTC	TAT	GTG	AGA	AAT	ATT	TGT	AAA	GCT	CTT	TTG	GAT	GTG	CTT	GGA	1152
Lys	Val	Tyr	Val	Arg	Asn	Ile	Cys	Lys	Ala	Leu	Leu	Asp	Val	Leu	Gly	
	370					375					380					
ATT	GAG	GTA	GAT	GCA	GAG	TAC	TTG	TTG	GGC	CCA	CTT	TAT	GCA	GCT	TTG	1200
Ile	Glu	Val	Asp	Ala	Glu	Tyr	Leu	Leu	Gly	Pro	Leu	Tyr	Ala	Ala	Leu	
385					390					395					400	
AAA	ATG	GAA	AGT	ATG	GAA	ATC	ATT	GAG	GAG	ATT	CAA	TGC	CAA	ACT	CAA	1248
Lys	Met	Glu	Ser	Met	Glu	Ile	Ile	Glu	Glu	Ile	Gln	Cys	Gln	Thr	Gln	
				405					410					415		
CAG	GAA	AAC	CTC	AGC	AGT	AAT	AGT	GAT	GGA	ATA	TCA	CCC	AAA	AGG	CGT	1296
Gln	Glu	Asn	Leu	Ser	Ser	Asn	Ser	Asp	Gly	Ile	Ser	Pro	Lys	Arg	Arg	
			420					425					430			
CGT	CTC	AGC	TCG	TCT	CTA	AAC	CCT	TCT	AAA	AGA	GCA	CCA	AAA	CAG	ACT	1344
Arg	Leu	Ser	Ser	Ser	Leu	Asn	Pro	Ser	Lys	Arg	Ala	Pro	Lys	Gln	Thr	
		435					440					445				
GAG	GAA	ATT	AAA	CAT	GTG	GAC	ATG	AAC	CAA	AAG	AGC	ATA	TTA	TGG	AGT	1392
Glu	Glu	Ile	Lys	His	Val	Asp	Met	Asn	Gln	Lys	Ser	Ile	Leu	Trp	Ser	
	450					455					460					
GCA	CTG	AAA	CAG	AAA	GCT	GAA	TCC	CTT	CAG	ATT	TCC	CTT	GAA	TAC	AGT	1440
Ala	Leu	Lys	Gln	Lys	Ala	Glu	Ser	Leu	Gln	Ile	Ser	Leu	Glu	Tyr	Ser	
465					470					475					480	
GGC	CTA	AAG	AAT	CCT	GTT	ATT	GAG	ATG	TTA	GAA	GGA	ATT	GCT	GTT	GTC	1488
Gly	Leu	Lys	Asn	Pro	Val	Ile	Glu	Met	Leu	Glu	Gly	Ile	Ala	Val	Val	
				485					490					495		
TTA	CAA	CTG	ACT	GCT	CTG	TGT	ACT	GTT	CAT	TGT	TCT	CAT	CAA	AAC	ATG	1536
Leu	Gln	Leu	Thr	Ala	Leu	Cys	Thr	Val	His	Cys	Ser	His	Gln	Asn	Met	
			500					505					510			
AAC	TGC	CGT	ACT	TTC	AAG	GAC	TGT	CAA	CAT	AAA	TCC	AAG	AAG	AAA	CCT	1584
Asn	Cys	Arg	Thr	Phe	Lys	Asp	Cys	Gln	His	Lys	Ser	Lys	Lys	Lys	Pro	
		515					520					525				
TCT	GTA	GTG	ATA	ACT	TGG	ATG	TCA	TTG	GAT	TTT	TAC	ACA	AAA	GTG	CTT	1632
Ser	Val	Val	Ile	Thr	Trp	Met	Ser	Leu	Asp	Phe	Tyr	Thr	Lys	Val	Leu	
	530					535					540					
AAG	AGC	TGT	AGA	AGT	TTG	TTA	GAA	TCT	GTT	CAG	AAA	CTG	GAC	CTG	GAG	1680
Lys	Ser	Cys	Arg	Ser	Leu	Leu	Glu	Ser	Val	Gln	Lys	Leu	Asp	Leu	Glu	
545					550					555					560	
GCA	ACC	ATT	GAT	AAG	GTG	GTG	AAA	ATT	TAT	GAT	GCT	TTG	ATT	TAT	ATG	1728
Ala	Thr	Ile	Asp	Lys	Val	Val	Lys	Ile	Tyr	Asp	Ala	Leu	Ile	Tyr	Met	
				565					570					575		

-133-

CAA	GTA	AAC	AGT	TCA	TTT	GAA	GAT	CAT	ATC	CTG	GAA	GAT	TTA	TGT	GGT	1776
Gln	Val	Asn	Ser	Ser	Phe	Glu	Asp	His	Ile	Leu	Glu	Asp	Leu	Cys	Gly	
			580					585					590			
ATG	CTC	TCA	CTT	CCA	TGG	ATT	TAT	TCC	CAT	TCT	GAT	GAT	GGC	TGT	TTA	1824
Met	Leu	Ser	Leu	Pro	Trp	Ile	Tyr	Ser	His	Ser	Asp	Asp	Gly	Cys	Leu	
		595					600					605				
AAG	TTG	ACC	ACA	TTT	GCC	GCT	AAT	CTT	CTA	ACA	TTA	AGC	TGT	AGG	ATT	1872
Lys	Leu	Thr	Thr	Phe	Ala		Asn	Leu	Leu	Thr		Ser	Cys	Arg	Ile	
	610					615					620					
TCA	GAT	AGC	TAT	TCA	CCA	CAG	GCA	CAA	TCA	CGA	TGT	GTG	TTT	CTT	CTG	1920
Ser	Asp	Ser	Tyr	Ser	Pro	Gln	Ala	Gln	Ser	Arg	Cys	Val	Phe	Leu	Leu	
	625				630					635					640	
ACT	CTG	TTT	CCA	AGA	AGA	ATA	TTC	CTT	GAG	TGG	AGA	ACA	GCA	GTT	TAC	1968
Thr	Leu	Phe	Pro	Arg	Arg	Ile	Phe	Leu	Glu	Trp	Arg	Thr	Ala	Val	Tyr	
			645						650					655		
AAC	TGG	GCC	CTG	CAG	AGC	TCC	CAT	GAA	GTA	ATC	CGG	GCT	AGT	TGT	GTT	2016
Asn	Trp	Ala	Leu	Gln	Ser	Ser	His	Glu	Val	Ile	Arg	Ala	Ser	Cys	Val	
		660						665					670			
AGT	GGA	TTT	TTT	ATC	TTA	TTG	CAG	CAG	CAG	AAT	TCT	TGT	AAC	AGA	GTT	2064
Ser	Gly	Phe	Phe	Ile	Leu	Leu	Gln	Gln	Gln	Asn	Ser	Cys	Asn	Arg	Val	
		675					680					685				
CCC	AAG	ATT	CTT	ATA	GAT	AAA	GTC	AAA	GAT	GAT	TCT	GAC	ATT	GTC	AAG	2112
Pro	Lys	Ile	Leu	Ile	Asp	Lys	Val	Lys	Asp	Asp	Ser	Asp	Ile	Val	Lys	
	690					695					700					
AAA	GAA	TTT	GCT	TCT	ATA	CTT	GGT	CAA	CTT	GTC	TGT	ACT	CTT	CAC	GGC	2160
Lys	Glu	Phe	Ala	Ser	Ile	Leu	Gly	Gln	Leu	Val	Cys	Thr	Leu	His	Gly	
	705				710					715					720	
ATG	TTT	TAT	CTG	ACA	AGT	TCT	TTA	ACA	GAA	CCT	TTC	TCT	GAA	CAC	GGA	2208
Met	Phe	Tyr	Leu	Thr	Ser	Ser	Leu	Thr	Glu	Pro	Phe	Ser	Glu	His	Gly	
				725					730					735		
CAT	GTG	GAC	CTC	TTC	TGT	AGG	AAC	TTG	AAA	GCC	ACT	TCT	CAA	CAT	GAA	2256
His	Val	Asp	Leu	Phe	Cys	Arg	Asn	Leu	Lys	Ala	Thr	Ser	Gln	His	Glu	
			740					745					750			
TGT	TCA	TCT	TCT	CAA	CTA	AAA	GCT	TCT	GTC	TGC	AAG	CCA	TTC	CTT	TTC	2304
Cys	Ser		Ser	Gln	Leu	Lys	Ala	Ser	Val	Cys	Lys	Pro	Phe	Leu	Phe	
		755					760					765				
CTA	CTG	AAA	AAA	AAA	ATA	CCT	AGT	CCA	GTA	AAA	CTT	GCT	TTC	ATA	GAT	2352
Leu	Leu	Lys	Lys	Lys	Ile	Pro	Ser	Pro	Val	Lys	Leu	Ala	Phe	Ile	Asp	
	770					775					780					
AAT	CTA	CAT	CAT	CTT	TGT	AAG	CAT	CTT	GAT	TTT	AGA	GAA	GAT	GAA	ACA	2400
Asn	Leu	His	His	Leu	Cys	Lys	His	Leu	Asp	Phe	Arg	Glu	Asp	Glu	Thr	
	785				790					795					800	
GAT	GTA	AAA	GCA	GTT	CTT	GGA	ACT	TTA	TTA	AAT	TTA	ATG	GAA	GAT	CCA	2448
Asp	Val	Lys	Ala	Val	Leu	Gly	Thr	Leu	Leu	Asn	Leu	Met	Glu	Asp	Pro	
				805				810					815			
GAC	AAA	GAT	GTT	AGA	GTG	GCT	TTT	AGT	GGA	AAT	ATC	AAG	CAC	ATA	TTG	2496
Asp	Lys	Asp	Val	Arg	Val	Ala	Phe	Ser	Gly	Asn	Ile	Lys	His	Ile	Leu	
			820					825					830			
GAA	TCC	TTG	GAC	TCT	GAA	GAT	GGA	TTT	ATA	AAG	GAG	CTT	TTT	GTC	TTA	2544
Glu	Ser	Leu	Asp	Ser	Glu	Asp	Gly	Phe	Ile	Lys	Glu	Leu	Phe	Val	Leu	
		835					840					845				

-134-

AGA	ATG	AAG	GAA	GCA	TAT	ACA	CAT	GCC	CAA	ATA	TCA	AGA	AAT	AAT	GAG	2592
Arg	Met	Lys	Glu	Ala	Tyr	Thr	His	Ala	Gln	Ile	Ser	Arg	Asn	Asn	Glu	
850						855					860					
CTG	AAG	GAT	ACC	TTG	ATT	CTT	ACA	ACA	GGG	GAT	ATT	GGA	AGG	GCC	GCA	2640
Leu	Lys	Asp	Thr	Leu	Ile	Leu	Thr	Thr	Gly	Asp	Ile	Gly	Arg	Ala	Ala	
865					870				875						880	
AAA	GGA	GAT	TTG	GTA	CCA	TTT	GCA	CTC	TTA	CAC	TTA	TTG	CAT	TGT	TTG	2688
Lys	Gly	Asp	Leu	Val	Pro	Phe	Ala	Leu	Leu	His	Leu	Leu	His	Cys	Leu	
				885					890					895		
TTA	TCC	AAG	TCA	GCA	TCT	GTC	TCT	GGA	GCA	GCA	TAC	ACA	GAA	ATT	AGA	2736
Leu	Ser	Lys	Ser	Ala	Ser	Val	Ser	Gly	Ala	Ala	Tyr	Thr	Glu	Ile	Arg	
			900					905					910			
GCT	CTG	GTT	GCA	GCT	AAA	AGT	GTT	AAA	CTG	CAA	AGT	TTT	TTC	AGC	CAG	2784
Ala	Leu	Val	Ala	Ala	Lys	Ser	Val	Lys	Leu	Gln	Ser	Phe	Phe	Ser	Gln	
		915					920					925				
TAT	AAG	AAA	CCC	ATC	TGT	CAG	TTT	TTG	GTA	GAA	TCC	CTT	CAC	TCT	AGT	2832
Tyr	Lys	Lys	Pro	Ile	Cys	Gln	Phe	Leu	Val	Glu	Ser	Leu	His	Ser	Ser	
	930					935					940					
CAG	ATG	ACA	GCA	CTT	CCG	AAT	ACT	CCA	TGC	CAG	AAT	GCT	GAC	GTG	CGA	2880
Gln	Met	Thr	Ala	Leu	Pro	Asn	Thr	Pro	Cys	Gln	Asn	Ala	Asp	Val	Arg	
945					950					955					960	
AAA	CAA	GAT	GTG	GCT	CAC	CAG	AGA	GAA	ATG	GCT	TTA	AAT	ACG	TTG	TCT	2928
Lys	Gln	Asp	Val	Ala	His	Gln	Arg	Glu	Met	Ala	Leu	Asn	Thr	Leu	Ser	
				965					970					975		
GAA	ATT	GCC	AAC	GTT	TTC	GAC	TTT	CCT	GAT	CTT	AAT	CGT	TTT	CTT	ACT	2976
Glu	Ile	Ala	Asn	Val	Phe	Asp	Phe	Pro	Asp	Leu	Asn	Arg	Phe	Leu	Thr	
			980					985					990			
AGG	ACA	TTA	CAA	GTT	CTA	CTA	CCT	GAT	CTT	GCT	GCC	AAA	GCA	AGC	CCT	3024
Arg	Thr	Leu	Gln	Val	Leu	Leu	Pro	Asp	Leu	Ala	Ala	Lys	Ala	Ser	Pro	
		995					1000					1005				
GCA	GCT	TCT	GCT	CTC	ATT	CGA	ACT	TTA	GGA	AAA	CAA	TTA	AAT	GTC	AAT	3072
Ala	Ala	Ser	Ala	Leu	Ile	Arg	Thr	Leu	Gly	Lys	Gln	Leu	Asn	Val	Asn	
	1010					1015					1020					
CGT	AGA	GAG	ATT	TTA	ATA	AAC	AAC	TTC	AAA	TAT	ATT	TTT	TCT	CAT	TTG	3120
Arg	Arg	Glu	Ile	Leu	Ile	Asn	Asn	Phe	Lys	Tyr	Ile	Phe	Ser	His	Leu	
1025					1030					1035					1040	
GTC	TGT	TCT	TGT	TCC	AAA	GAT	GAA	TTA	GAA	CGT	GCC	CTT	CAT	TAT	CTG	3168
Val	Cys	Ser	Cys	Ser	Lys	Asp	Glu	Leu	Glu	Arg	Ala	Leu	His	Tyr	Leu	
				1045					1050					1055		
AAG	AAT	GAA	ACA	GAA	ATT	GAA	CTG	GGG	AGC	CTG	TTG	AGA	CAA	GAT	TTC	3216
Lys	Asn	Glu	Thr	Glu	Ile	Glu	Leu	Gly	Ser	Leu	Leu	Arg	Gln	Asp	Phe	
			1060					1065					1070			
CAA	GGA	TTG	CAT	AAT	GAA	TTA	TTG	CTG	CGT	ATT	GGA	GAA	CAC	TAT	CAA	3264
Gln	Gly	Leu	His	Asn	Glu	Leu	Leu	Leu	Arg	Ile	Gly	Glu	His	Tyr	Gln	
		1075					1080					1085				
CAG	GTT	TTT	AAT	GGT	TTG	TCA	ATA	CTT	GCC	TCA	TTT	GCA	TCC	AGT	GAT	3312
Gln	Val	Phe	Asn	Gly	Leu	Ser	Ile	Leu	Ala	Ser	Phe	Ala	Ser	Ser	Asp	
	1090					1095					1100					
GAT	CCA	TAT	CAG	GGC	CCG	AGA	GAT	ATC	ATA	TCA	CCT	GAA	CTG	ATG	GCT	3360
Asp	Pro	Tyr	Gln	Gly	Pro	Arg	Asp	Ile	Ile	Ser	Pro	Glu	Leu	Met	Ala	
1105					1110					1115					1120	

-135-

GAT TAT TTA CAA CCC AAA TTG TTG GGC ATT TTG GCT TTT TTT AAC ATG Asp Tyr Leu Gln Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met 1125 1130 1135	3408
CAG TTA CTG AGC TCT AGT GTT GGC ATT GAA GAT AAG AAA ATG GCC TTG Gln Leu Leu Ser Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu 1140 1145 1150	3456
AAC AGT TTG ATG TCT TTG ATG AAG TTA ATG GGA CCC AAA CAT GTC AGT Asn Ser Leu Met Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser 1155 1160 1165	3504
TCT GTG AGG GTG AAG ATG ATG ACC ACA CTG AGA ACT GGC CTT CGA TTC Ser Val Arg Val Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe 1170 1175 1180	3552
AAG GAT GAT TTT CCT GAA TTG TGT TGC AGA GCT TGG GAC TGC TTT GTT Lys Asp Asp Phe Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val 1185 1190 1195 1200	3600
CGC TGC CTG GAT CAT GCT TGT CTG GGC TCC CTT CTC AGT CAT GTA ATA Arg Cys Leu Asp His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile 1205 1210 1215	3648
GTA GCT TTG TTA CCT CTT ATA CAC ATC CAG CCT AAA GAA ACT GCA GCT Val Ala Leu Leu Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala 1220 1225 1230	3696
ATC TTC CAC TAC CTC ATA ATT GAA AAC AGG GAT GCT GTG CAA GAT TTT Ile Phe His Tyr Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe 1235 1240 1245	3744
CTT CAT GAA ATA TAT TTT TTA CCT GAT CAT CCA GAA TTA AAA AAG ATA Leu His Glu Ile Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile 1250 1255 1260	3792
AAA GCC GTT CTC CAG GAA TAC AGA AAG GAG ACC TCT GAG AGC ACT GAT Lys Ala Val Leu Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp 1265 1270 1275 1280	3840
CTT CAG ACA ACT CTT CAG CTC TCT ATG AAG GCC ATT CAA CAT GAA AAT Leu Gln Thr Thr Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn 1285 1290 1295	3888
GTC GAT GTT CGT ATT CAT GCT CTT ACA AGC TTG AAG GAA ACC TTG TAT Val Asp Val Arg Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr 1300 1305 1310	3936
AAA AAT CAG GAA AAA CTG ATA AAG TAT GCA ACA GAC AGT GAA ACA GTA Lys Asn Gln Glu Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val 1315 1320 1325	3984
GAA CCT ATT ATC TCA CAG TTG GTG ACA GTG CTT TTG AAA GGT TGC CAA Glu Pro Ile Ile Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln 1330 1335 1340	4032
GAT GCA AAC TCT CAA GCT CGG TTG CTC TGT GGG GAA TGT TTA GGG GAA Asp Ala Asn Ser Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu 1345 1350 1355 1360	4080
TTG GGG GCG ATA GAT CCA GGT CGA TTA GAT TTC TCA ACA ACT GAA ACT Leu Gly Ala Ile Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr 1365 1370 1375	4128
CAA GGA AAA GAT TTT ACA TTT GTG ACT GGA GTA GAA GAT TCA AGC TTT Gln Gly Lys Asp Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe 1380 1385 1390	4176

-136-

GCC TAT GGA TTA TTG ATG GAG CTA ACA AGA GCT TAC CTT GCG TAT GCT Ala Tyr Gly Leu Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala 1395 1400 1405	4224
GAT AAT AGC CGA GCT CAA GAT TCA GCT GCC TAT GCC ATT CAG GAG TTG Asp Asn Ser Arg Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu 1410 1415 1420	4272
CTT TCT ATT TAT GAC TGT AGA GAG ATG GAG ACC AAC GGC CCA GGT CAC Leu Ser Ile Tyr Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His 1425 1430 1435 1440	4320
CAA TTG TGG AGG AGA TTT CCT GAG CAT GTT CGG GAA ATA CTA GAA CCT Gln Leu Trp Arg Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro 1445 1450 1455	4368
CAT CTA AAT ACC AGA TAC AAG AGT TCT CAG AAG TCA ACC GAT TGG TCT His Leu Asn Thr Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser 1460 1465 1470	4416
GGA GTA AAG AAG CCA ATT TAC TTA AGT AAA TTG GGT AGT AAC TTT GCA Gly Val Lys Lys Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala 1475 1480 1485	4464
GAA TGG TCA GCA TCT TGG GCA GGT TAT CTT ATT ACA AAG GTT CGA CAT Glu Trp Ser Ala Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His 1490 1495 1500	4512
GAT CTT GCC AGT AAA ATT TTC ACC TGC TGT AGC ATT ATG ATG AAG CAT Asp Leu Ala Ser Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His 1505 1510 1515 1520	4560
GAT TTC AAA GTG ACC ATC TAT CTT CTT CCA CAT ATT CTG GTG TAT GTC Asp Phe Lys Val Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val 1525 1530 1535	4608
TTA CTG GGT TGT AAT CAA GAA GAT CAG CAG GAG GTT TAT GCA GAA ATT Leu Leu Gly Cys Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile 1540 1545 1550	4656
ATG GCA GTT CTA AAG CAT GAC GAT CAG CAT ACC ATA AAT ACC CAA GAC Met Ala Val Leu Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp 1555 1560 1565	4704
ATT GCA TCT GAT CTG TGT CAA CTC AGT ACA CAG ACT GTG TTC TCC ATG Ile Ala Ser Asp Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met 1570 1575 1580	4752
CTT GAC CAT CTC ACA CAG TGG GCA AGG CAC AAA TTT CAG GCA CTG AAA Leu Asp His Leu Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys 1585 1590 1595 1600	4800
GCT GAG AAA TGT CCA CAC AGC AAA TCA AAC AGA AAT AAG GTA GAC TCA Ala Glu Lys Cys Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser 1605 1610 1615	4848
ATG GTA TCT ACT GTG GAT TAT GAA GAC TAT CAG AGT GTA ACC CGT TTT Met Val Ser Thr Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe 1620 1625 1630	4896
CTA GAC CTC ATA CCC CAG GAT ACT CTG GCA GTA GCT TCC TTT CGC TCC Leu Asp Leu Ile Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser 1635 1640 1645	4944
AAA GCA TAC ACA CGA GCT GTA ATG CAC TTT GAA TCA TTT ATT ACA GAA Lys Ala Tyr Thr Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu 1650 1655 1660	4992

-137-

AAG Lys 1665	AAG Lys 1665	CAA Gln 1665	AAT Asn 1665	ATT Ile 1665	CAG Gln 1670	GAA Glu 1670	CAT His 1670	CTT Leu 1670	GGA Gly 1675	TTT Phe 1675	TTA Leu 1675	CAG Gln 1675	AAA Lys 1675	TTG Leu 1680	TAT Tyr 1680	5040
GCT Ala 1685	GCT Ala 1685	ATG Met 1685	CAT His 1685	GAA Glu 1685	CCT Pro 1685	GAT Asp 1685	GGA Gly 1690	GTG Val 1690	GCC Ala 1690	GGA Gly 1690	GTC Val 1690	AGT Ser 1695	GCA Ala 1695	ATT Ile 1695	AGA Arg 1695	5088
AAG Lys 1700	GCA Ala 1700	GAA Glu 1700	CCA Pro 1700	TCT Ser 1700	CTA Leu 1705	AAA Lys 1705	GAA Glu 1705	CAG Gln 1705	ATC Ile 1705	CTT Leu 1710	GAA Glu 1710	CAT His 1710	GAA Glu 1710	AGC Ser 1710	CTT Leu 1710	5136
GGC Gly 1715	TTG Leu 1715	CTG Leu 1715	AGG Arg 1715	GAT Asp 1720	GCC Ala 1720	ACT Thr 1720	GCT Ala 1720	TGT Cys 1720	TAT Tyr 1725	GAC Asp 1725	AGG Arg 1725	GCT Ala 1725	ATT Ile 1725	CAG Gln 1725	CTA Leu 1725	5184
GAA Glu 1730	CCA Pro 1730	GAC Asp 1730	CAG Gln 1735	ATC Ile 1735	ATT Ile 1735	CAT His 1735	TAT Tyr 1735	CAT His 1740	GGT Gly 1740	GTA Val 1740	GTA Val 1740	AAG Lys 1740	TCC Ser 1740	ATG Met 1740	TTA Leu 1740	5232
GGT Gly 1745	CTT Leu 1745	GGT Gly 1750	CAG Gln 1750	CTG Leu 1750	TCT Ser 1750	ACT Thr 1755	GTT Val 1755	ATC Ile 1755	ACT Thr 1755	CAG Gln 1755	GTG Val 1755	AAT Asn 1760	GGA Gly 1760	GTG Val 1760	CAT His 1760	5280
GCT Ala 1765	AAC Asn 1765	AGG Arg 1765	TCC Ser 1765	GAG Glu 1765	TGG Trp 1765	ACA Thr 1770	GAT Asp 1770	GAA Glu 1770	TTA Leu 1770	AAC Asn 1775	ACG Thr 1775	TAC Tyr 1775	AGA Arg 1775	GTG Val 1775	GAA Glu 1775	5328
GCA Ala 1780	GCT Ala 1780	TGG Trp 1780	AAA Lys 1780	TTG Leu 1780	TCA Ser 1785	CAG Gln 1785	TGG Trp 1785	GAT Asp 1785	TTG Leu 1785	GTG Val 1790	GAA Glu 1790	AAC Asn 1790	TAT Tyr 1790	TTG Leu 1790	GCA Ala 1790	5376
GCA Ala 1795	GAT Asp 1795	GGA Gly 1795	AAA Lys 1795	TCT Ser 1795	ACA Thr 1800	ACA Thr 1800	TGG Trp 1800	AGT Ser 1800	GTC Val 1805	AGA Arg 1805	CTG Leu 1805	GGA Gly 1805	CAG Gln 1805	CTA Leu 1805	TTA Leu 1805	5424
TTA Leu 1810	TCA Ser 1810	GCC Ala 1810	AAA Lys 1815	AAA Lys 1815	AGA Arg 1815	GAT Asp 1815	ATC Ile 1815	ACA Thr 1820	GCT Ala 1820	TTT Phe 1820	TAT Tyr 1820	GAC Asp 1820	TCA Ser 1820	CTG Leu 1820	AAA Lys 1820	5472
CTA Leu 1825	GTG Val 1825	AGA Arg 1830	GCA Ala 1830	GAA Glu 1830	CAA Gln 1830	ATT Ile 1835	GTA Val 1835	CCT Pro 1835	CTT Leu 1835	TCA Ser 1835	GCT Ala 1835	GCA Ala 1835	AGC Ser 1840	TTT Phe 1840	GAA Glu 1840	5520
AGA Arg 1845	GGC Gly 1845	TCC Ser 1845	TAC Tyr 1845	CAA Gln 1845	CGA Arg 1850	GGA Gly 1850	TAT Tyr 1850	GAA Glu 1850	TAT Tyr 1850	ATT Ile 1855	GTG Val 1855	AGA Arg 1855	TTG Leu 1855	CAC His 1855	ATG Met 1855	5568
TTA Leu 1860	TGT Cys 1860	GAG Glu 1860	TTG Leu 1860	GAG Glu 1860	CAT His 1865	AGC Ser 1865	ATC Ile 1865	AAA Lys 1865	CCA Pro 1865	CTT Leu 1865	TTC Phe 1870	CAG Gln 1870	CAT His 1870	TCT Ser 1870	CCA Pro 1870	5616
GGT Gly 1875	GAC Asp 1875	AGT Ser 1875	TCT Ser 1875	CAA Gln 1880	GAA Glu 1880	GAT Asp 1880	TCT Ser 1880	CTA Leu 1880	AAC Asn 1885	TGG Trp 1885	GTA Val 1885	GCT Ala 1885	CGA Arg 1885	CTA Leu 1885	GAA Glu 1885	5664
ATG Met 1890	ACC Thr 1890	CAG Gln 1895	AAT Asn 1895	TCC Ser 1895	TAC Tyr 1895	AGA Arg 1895	GCC Ala 1895	AAG Lys 1900	GAG Glu 1900	CCT Pro 1900	ATC Ile 1900	CTG Leu 1900	GCT Ala 1900	CTC Leu 1900	CGG Arg 1900	5712
AGG Arg 1905	GCT Ala 1905	TTA Leu 1905	CTA Leu 1905	AGC Ser 1910	CTC Leu 1910	AAC Asn 1910	AAA Lys 1915	AGA Arg 1915	CCA Pro 1915	GAT Asp 1915	TAC Tyr 1915	AAT Asn 1915	GAA Glu 1920	ATG Met 1920	GTT Val 1920	5760
GGA Gly 1925	GAA Glu 1925	TGC Cys 1925	TGG Trp 1925	CTG Leu 1925	CAG Gln 1930	AGT Ser 1930	GCC Ala 1930	AGG Arg 1930	GTA Val 1930	GCT Ala 1930	AGA Arg 1930	AAG Lys 1930	GCT Ala 1930	GGT Gly 1935	CAC His 1935	5808

-138-

CAC CAG ACA GCC TAC AAT GCT CTC CTT AAT GCA GGG GAA TCA CGA CTC His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu 1940 1945 1950	5856
GCT GAA CTG TAC GTG GAA AGG GCA AAG TGG CTC TGG TCC AAG GGT GAT Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp 1955 1960 1965	5904
GTT CAC CAG GCA CTA ATT GTT CTT CAA AAA GGT GTT GAA TTA TGT TTT Val His Gln Ala Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe 1970 1975 1980	5952
CCT GAA AAT GAA ACC CCA CCT GAG GGT AAG AAC ATG TTA ATC CAT GGT Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly 1985 1990 1995 2000	6000
CGA GCT ATG CTA CTA GTG GGC CGA TTT ATG GAA GAA ACA GCT AAC TTT Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe 2005 2010 2015	6048
GAA AGC AAT GCA ATT ATG AAA AAA TAT AAG GAT GTG ACC GCG TGC CTG Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu 2020 2025 2030	6096
CCA GAA TGG GAG GAT GGG CAT TTT TAC CTT GCC AAG TAC TAT GAC AAA Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys 2035 2040 2045	6144
TTG ATG CCC ATG GTC ACA GAC AAC AAA ATG GAA AAG CAA GGT GAT CTC Leu Met Pro Met Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu 2050 2055 2060	6192
ATC CGG TAT ATA GTT CTT CAT TTT GGC AGA TCT CTA CAA TAT GGA AAT Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn 2065 2070 2075 2080	6240
CAG TTC ATA TAT CAG TCA ATG CCA CGA ATG TTA ACT CTA TGG CTT GAT Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp 2085 2090 2095	6288
TAT GGT ACA AAG GCA TAT GAA TGG GAA AAA GCT GGC CGC TCC GAT CGT Tyr Gly Thr Lys Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg 2100 2105 2110	6336
GTA CAA ATG AGG AAT GAT TTG GGT AAA ATA AAC AAG GTT ATC ACA GAG Val Gln Met Arg Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu 2115 2120 2125	6384
CAT ACA AAC TAT TTA GCT CCA TAT CAA TTT TTG ACT GCT TTT TCA CAA His Thr Asn Tyr Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln 2130 2135 2140	6432
TTG ATC TCT CGA ATT TGT CAT TCT CAC GAT GAA GTT TTT GTT GTC TTG Leu Ile Ser Arg Ile Cys His Ser His Asp Glu Val Phe Val Val Leu 2145 2150 2155 2160	6480
ATG GAA ATA ATA GCC AAA GTA TTT CTA GCC TAT CCT CAA CAA GCA ATG Met Glu Ile Ile Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln Ala Met 2165 2170 2175	6528
TGG ATG ATG ACA GCT GTG TCA AAG TCA TCT TAT CCC ATG CGT GTG AAC Trp Met Met Thr Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn 2180 2185 2190	6576
AGA TGC AAG GAA ATC CTC AAT AAA GCT ATT CAT ATG AAA AAA TCC TTA Arg Cys Lys Glu Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu 2195 2200 2205	6624



-139-

GAG AAG TTT GTT GGA GAT GCA ACT CGC CTA ACA GAT AAG CTT CTA GAA Glu Lys Phe Val Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu 2210 2215 2220	6672
TTG TGC AAT AAA CCG GTT GAT GGA AGT AGT TCC ACA TTA AGC ATG AGC Leu Cys Asn Lys Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser 2225 2230 2235 2240	6720
ACT CAT TTT AAA ATG CTT AAA AAG CTG GTA GAA GAA GCA ACA TTT AGT Thr His Phe Lys Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser 2245 2250 2255	6768
GAA ATC CTC ATT CCT CTA CAA TCA GTC ATG ATA CCT ACA CTT CCA TCA Glu Ile Leu Ile Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser 2260 2265 2270	6816
ATT CTG GGT ACC CAT GCT AAC CAT GCT AGC CAT GAA CCA TTT CCT GGA Ile Leu Gly Thr His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly 2275 2280 2285	6864
CAT TGG GCC TAT ATT GCA GGG TTT GAT GAT ATG GTG GAA ATT CTT GCT His Trp Ala Tyr Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala 2290 2295 2300	6912
TCT CTT CAG AAA CCA AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys 2305 2310 2315 2320	6960
TTC TAC ATC ATG ATG TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT Phe Tyr Ile Met Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys 2325 2330 2335	7008
AGA CTA ATG GAA TTC AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT Arg Leu Met Glu Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp 2340 2345 2350	7056
GCA GAG TCT CGT AGA AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT Ala Glu Ser Arg Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile 2355 2360 2365	7104
CCA CTA AAT GAT GAA TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala 2370 2375 2380	7152
GGT TTG AGA CCT ATT CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr 2385 2390 2395 2400	7200
ATG ACA GGA AAA GAA CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT Met Thr Gly Lys Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala 2405 2410 2415	7248
TTA TCT GAA AAA CTC AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT Leu Ser Glu Lys Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His 2420 2425 2430	7296
CCT CCT ATT TTT CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA Pro Pro Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr 2435 2440 2445	7344
TCA TGG TAC AGT AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met 2450 2455 2460	7392
TCA ATG GTT GGT TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT Ser Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn 2465 2470 2475 2480	7440

-140-

ATT CTC TTT GAT TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn 2485 2490 2495	7488
TGT CTT TTC AAT AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro 2500 2505 2510	7536
TTT CGC CTG ACT CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA Phe Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr 2515 2520 2525	7584
GAG GGT CTT TTT CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT Glu Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg 2530 2535 2540	7632
GAT CAG CGA GAG CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT Asp Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp 2545 2550 2555 2560	7680
CCT CTT GTG GAA TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA Pro Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro 2565 2570 2575	7728
CTG AAT GAA ACT GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT Leu Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val 2580 2585 2590	7776
CTT GAC ATT GAG CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA Leu Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg 2595 2600 2605	7824
GTG ACA GGA CTG CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA Val Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile 2610 2615 2620	7872
CAA GAA GCT ACT GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp 2625 2630 2635 2640	7920
ACT CCA TAT ATG TGA Thr Pro Tyr Met	7935

## (2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2644 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

Met Gly Glu His Gly Leu Glu Leu Ala Ser Met Ile Pro Ala Leu Arg 1 5 10 15
Glu Leu Gly Ser Ala Thr Pro Glu Glu Tyr Asn Thr Val Val Gln Lys 20 25 30
Pro Arg Gln Ile Leu Cys Gln Phe Ile Asp Arg Ile Leu Thr Asp Val 35 40 45

-141-

Asn Val Val Ala Val Glu Leu Val Lys Lys Thr Asp Ser Gln Pro Thr  
 50 55 60  
 Ser Val Met Leu Leu Asp Phe Ile Gln His Ile Met Lys Ser Ser Pro  
 65 70 75 80  
 Leu Met Phe Val Asn Val Ser Gly Ser His Glu Arg Lys Gly Ser Cys  
 85 90 95  
 Ile Glu Phe Ser Asn Trp Ile Ile Thr Arg Leu Leu Arg Ile Ala Ala  
 100 105 110  
 Thr Pro Ser Cys His Leu Leu His Lys Lys Ile Cys Glu Val Ile Cys  
 115 120 125  
 Ser Leu Leu Phe Leu Phe Lys Ser Lys Ser Pro Ala Ile Phe Gly Val  
 130 135 140  
 Leu Thr Lys Glu Leu Leu Gln Leu Phe Glu Asp Leu Val Tyr Leu His  
 145 150 155 160  
 Arg Arg Asn Val Met Gly His Ala Val Glu Trp Pro Val Val Met Ser  
 165 170 175  
 Arg Phe Leu Ser Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala  
 180 185 190  
 Pro Leu Gln Leu Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr  
 195 200 205  
 Leu Leu Met Val Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg  
 210 215 220  
 Gln Glu Leu Leu Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly  
 225 230 235 240  
 Ser Pro Lys Ile Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe  
 245 250 255  
 Gln Leu Gly Gly Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser  
 260 265 270  
 Phe Leu Glu Leu Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu  
 275 280 285  
 Lys Leu Tyr Glu Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro  
 290 295 300  
 Phe Glu Ala Glu Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met  
 305 310 315 320  
 Leu Leu Glu Lys Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg  
 325 330 335  
 Leu Lys Ser Asp Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr  
 340 345 350  
 Phe Leu Lys Phe Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg  
 355 360 365  
 Lys Val Tyr Val Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly  
 370 375 380  
 Ile Glu Val Asp Ala Glu Tyr Leu Leu Gly Pro Leu Tyr Ala Ala Leu  
 385 390 395 400

-142-

Lys Met Glu Ser Met Glu Ile Ile Glu Glu Ile Gln Cys Gln Thr Gln  
 405 410 415  
 Gln Glu Asn Leu Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg  
 420 425 430  
 Arg Leu Ser Ser Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr  
 435 440 445  
 Glu Glu Ile Lys His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser  
 450 455 460  
 Ala Leu Lys Gln Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser  
 465 470 475 480  
 Gly Leu Lys Asn Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val  
 485 490 495  
 Leu Gln Leu Thr Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met  
 500 505 510  
 Asn Cys Arg Thr Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro  
 515 520 525  
 Ser Val Val Ile Thr Trp Met Ser Leu Asp Phe Tyr Thr Lys Val Leu  
 530 535 540  
 Lys Ser Cys Arg Ser Leu Leu Glu Ser Val Gln Lys Leu Asp Leu Glu  
 545 550 555 560  
 Ala Thr Ile Asp Lys Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met  
 565 570 575  
 Gln Val Asn Ser Ser Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly  
 580 585 590  
 Met Leu Ser Leu Pro Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu  
 595 600 605  
 Lys Leu Thr Thr Phe Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile  
 610 615 620  
 Ser Asp Ser Tyr Ser Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu  
 625 630 635 640  
 Thr Leu Phe Pro Arg Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr  
 645 650 655  
 Asn Trp Ala Leu Gln Ser Ser His Glu Val Ile Arg Ala Ser Cys Val  
 660 665 670  
 Ser Gly Phe Phe Ile Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val  
 675 680 685  
 Pro Lys Ile Leu Ile Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys  
 690 695 700  
 Lys Glu Phe Ala Ser Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly  
 705 710 715 720  
 Met Phe Tyr Leu Thr Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly  
 725 730 735  
 His Val Asp Leu Phe Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu  
 740 745 750

-143-

Cys Ser Ser Ser Gln Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe  
 755 760 765  
 Leu Leu Lys Lys Lys Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp  
 770 775 780  
 Asn Leu His His Leu Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr  
 785 790 795 800  
 Asp Val Lys Ala Val Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro  
 805 810 815  
 Asp Lys Asp Val Arg Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu  
 820 825 830  
 Glu Ser Leu Asp Ser Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu  
 835 840 845  
 Arg Met Lys Glu Ala Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu  
 850 855 860  
 Leu Lys Asp Thr Leu Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala  
 865 870 875 880  
 Lys Gly Asp Leu Val Pro Phe Ala Leu Leu His Leu Leu His Cys Leu  
 885 890 895  
 Leu Ser Lys Ser Ala Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg  
 900 905 910  
 Ala Leu Val Ala Ala Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln  
 915 920 925  
 Tyr Lys Lys Pro Ile Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser  
 930 935 940  
 Gln Met Thr Ala Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg  
 945 950 955 960  
 Lys Gln Asp Val Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser  
 965 970 975  
 Glu Ile Ala Asn Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr  
 980 985 990  
 Arg Thr Leu Gln Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro  
 995 1000 1005  
 Ala Ala Ser Ala Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn  
 1010 1015 1020  
 Arg Arg Glu Ile Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu  
 1025 1030 1035 1040  
 Val Cys Ser Cys Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu  
 1045 1050 1055  
 Lys Asn Glu Thr Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe  
 1060 1065 1070  
 Gln Gly Leu His Asn Glu Leu Leu Leu Arg Ile Gly Glu His Tyr Gln  
 1075 1080 1085  
 Gln Val Phe Asn Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp  
 1090 1095 1100

-144-

Asp Pro Tyr Gln Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala  
 1105 1110 1115 1120  
 Asp Tyr Leu Gln Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met  
 1125 1130 1135  
 Gln Leu Leu Ser Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu  
 1140 1145 1150  
 Asn Ser Leu Met Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser  
 1155 1160 1165  
 Ser Val Arg Val Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe  
 1170 1175 1180  
 Lys Asp Asp Phe Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val  
 1185 1190 1195 1200  
 Arg Cys Leu Asp His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile  
 1205 1210 1215  
 Val Ala Leu Leu Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala  
 1220 1225 1230  
 Ile Phe His Tyr Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe  
 1235 1240 1245  
 Leu His Glu Ile Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile  
 1250 1255 1260  
 Lys Ala Val Leu Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp  
 1265 1270 1275 1280  
 Leu Gln Thr Thr Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn  
 1285 1290 1295  
 Val Asp Val Arg Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr  
 1300 1305 1310  
 Lys Asn Gln Glu Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val  
 1315 1320 1325  
 Glu Pro Ile Ile Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln  
 1330 1335 1340  
 Asp Ala Asn Ser Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu  
 1345 1350 1355 1360  
 Leu Gly Ala Ile Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr  
 1365 1370 1375  
 Gln Gly Lys Asp Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe  
 1380 1385 1390  
 Ala Tyr Gly Leu Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala  
 1395 1400 1405  
 Asp Asn Ser Arg Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu  
 1410 1415 1420  
 Leu Ser Ile Tyr Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His  
 1425 1430 1435 1440  
 Gln Leu Trp Arg Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro  
 1445 1450 1455

-145-

His Leu Asn Thr Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser  
 1460 1465 1470  
 Gly Val Lys Lys Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala  
 1475 1480 1485  
 Glu Trp Ser Ala Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His  
 1490 1495 1500  
 Asp Leu Ala Ser Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His  
 1505 1510 1515 1520  
 Asp Phe Lys Val Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val  
 1525 1530 1535  
 Leu Leu Gly Cys Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile  
 1540 1545 1550  
 Met Ala Val Leu Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp  
 1555 1560 1565  
 Ile Ala Ser Asp Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met  
 1570 1575 1580  
 Leu Asp His Leu Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys  
 1585 1590 1595 1600  
 Ala Glu Lys Cys Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser  
 1605 1610 1615  
 Met Val Ser Thr Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe  
 1620 1625 1630  
 Leu Asp Leu Ile Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser  
 1635 1640 1645  
 Lys Ala Tyr Thr Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu  
 1650 1655 1660  
 Lys Lys Gln Asn Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr  
 1665 1670 1675 1680  
 Ala Ala Met His Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg  
 1685 1690 1695  
 Lys Ala Glu Pro Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu  
 1700 1705 1710  
 Gly Leu Leu Arg Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu  
 1715 1720 1725  
 Glu Pro Asp Gln Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu  
 1730 1735 1740  
 Gly Leu Gly Gln Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His  
 1745 1750 1755 1760  
 Ala Asn Arg Ser Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu  
 1765 1770 1775  
 Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala  
 1780 1785 1790  
 Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu  
 1795 1800 1805

-146-

Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys  
 1810 1815 1820  
 Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu  
 1825 1830 1835 1840  
 Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met  
 1845 1850 1855  
 Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro  
 1860 1865 1870  
 Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu  
 1875 1880 1885  
 Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg  
 1890 1895 1900  
 Arg Ala Leu Leu Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val  
 1905 1910 1915 1920  
 Gly Glu Cys Trp Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His  
 1925 1930 1935  
 His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu  
 1940 1945 1950  
 Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp  
 1955 1960 1965  
 Val His Gln Ala Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe  
 1970 1975 1980  
 Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly  
 1985 1990 1995 2000  
 Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe  
 2005 2010 2015  
 Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu  
 2020 2025 2030  
 Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys  
 2035 2040 2045  
 Leu Met Pro Met Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu  
 2050 2055 2060  
 Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn  
 2065 2070 2075 2080  
 Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp  
 2085 2090 2095  
 Tyr Gly Thr Lys Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg  
 2100 2105 2110  
 Val Gln Met Arg Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu  
 2115 2120 2125  
 His Thr Asn Tyr Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln  
 2130 2135 2140  
 Leu Ile Ser Arg Ile Cys His Ser His Asp Glu Val Phe Val Val Leu  
 2145 2150 2155 2160



-147-

Met Glu Ile Ile Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln Ala Met  
2165 2170 2175

Trp Met Met Thr Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn  
2180 2185 2190

Arg Cys Lys Glu Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu  
2195 2200 2205

Glu Lys Phe Val Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu  
2210 2215 2220

Leu Cys Asn Lys Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser  
2225 2230 2235 2240

Thr His Phe Lys Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser  
2245 2250 2255

Glu Ile Leu Ile Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser  
2260 2265 2270

Ile Leu Gly Thr His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly  
2275 2280 2285

His Trp Ala Tyr Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala  
2290 2295 2300

Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys  
2305 2310 2315 2320

Phe Tyr Ile Met Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys  
2325 2330 2335

Arg Leu Met Glu Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp  
2340 2345 2350

Ala Glu Ser Arg Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile  
2355 2360 2365

Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala  
2370 2375 2380

Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr  
2385 2390 2395 2400

Met Thr Gly Lys Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala  
2405 2410 2415

Leu Ser Glu Lys Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His  
2420 2425 2430

Pro Pro Ile Phe His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr  
2435 2440 2445

Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met  
2450 2455 2460

Ser Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn  
2465 2470 2475 2480

Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn  
2485 2490 2495

Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro  
2500 2505 2510

-148-

Phe Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr  
 2515 2520 2525  
 Glu Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg  
 2530 2535 2540  
 Asp Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp  
 2545 2550 2555 2560  
 Pro Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro  
 2565 2570 2575  
 Leu Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val  
 2580 2585 2590  
 Leu Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg  
 2595 2600 2605  
 Val Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile  
 2610 2615 2620  
 Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp  
 2625 2630 2635 2640  
 Thr Pro Tyr Met

## (2) INFORMATION FOR SEQ ID NO:30:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7624 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 333..7562

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

CTTGTGAAGA GAATGTTTTA CACTCTTGTT AGTGAAGTTT ATTCTTTAAA AGTCAATCGT 60  
 CAAGGATTTA GCAAATGAAT TAGCACTTCG GATATACTTG TTTATTTAAT ATCTTTTTTG 120  
 TTTATTTCAA AGAATTCAGT AATTGGATCA TAACGAGACT TCTGCGGATT GCAGCAACTC 180  
 CCTCCTGTCA TTTGTTACAC AAGAAAATCT GTGAAGTCAT CTGTTTATTA TTATTTCTTT 240  
 TTAAAAGCAA GAGTCCTGCT ATTTTGGGG TACTCACAAA AGAATTATTA CAACTTTTTG 300  
 AAGACTTGGT TTACCTCCAT AGAAGAAATG TG ATG GGT CAT GCT GTG GAA TGG 353  
 Met Gly His Ala Val Glu Trp  
 1 5  
 CCA GTG GTC ATG AGC CGA TTT TTA AGT CAA TTA GAT GAA CAC ATG GGA 401  
 Pro Val Val Met Ser Arg Phe Leu Ser Gln Leu Asp Glu His Met Gly  
 10 15 20  
 TAT TTA CAA TCA GCT CCT TTG CAG TTG ATG AGT ATG CAA AAT TTA GAA 449  
 Tyr Leu Gln Ser Ala Pro Leu Gln Leu Met Ser Met Gln Asn Leu Glu  
 25 30 35

-149-

TTT	ATT	GAA	GTC	ACT	TTA	TTA	ATG	GTT	CTT	ACT	CGT	ATT	ATT	GCA	ATT	497
Phe	Ile	Glu	Val	Thr	Leu	Leu	Met	Val	Leu	Thr	Arg	Ile	Ile	Ala	Ile	
40					45					50					55	
GTG	TTT	TTT	AGA	AGG	CAA	GAA	CTC	TTA	CTT	TGG	CAG	ATA	GGT	TGT	GTT	545
Val	Phe	Phe	Arg	Arg	Gln	Glu	Leu	Leu	Leu	Trp	Gln	Ile	Gly	Cys	Val	
				60					65					70		
CTG	CTA	GAG	TAT	GGT	AGT	CCA	AAA	ATT	AAA	TCC	CTA	GCA	ATT	AGC	TTT	593
Leu	Leu	Glu	Tyr	Gly	Ser	Pro	Lys	Ile	Lys	Ser	Leu	Ala	Ile	Ser	Phe	
			75					80					85			
TTA	ACA	GAA	CTT	TTT	CAG	CTT	GGA	GGA	CTA	CCA	GCA	CAA	CCA	GCT	AGC	641
Leu	Thr	Glu	Leu	Phe	Gln	Leu	Gly	Gly	Leu	Pro	Ala	Gln	Pro	Ala	Ser	
		90					95					100				
ACT	TTT	TTC	AGC	TCA	TTT	TTG	GAA	TTA	TTA	AAA	CAC	CTT	GTA	GAA	ATG	689
Thr	Phe	Phe	Ser	Ser	Phe	Leu	Glu	Leu	Leu	Lys	His	Leu	Val	Glu	Met	
	105					110					115					
GAT	ACT	GAC	CAA	TTG	AAA	CTC	TAT	GAA	GAG	CCA	TTA	TCA	AAG	CTG	ATA	737
Asp	Thr	Asp	Gln	Leu	Lys	Leu	Tyr	Glu	Glu	Pro	Leu	Ser	Lys	Leu	Ile	
120					125					130					135	
AAG	ACA	CTA	TTT	CCC	TTT	GAA	GCA	GAA	GCT	TAT	AGA	AAT	ATT	GAA	CCT	785
Lys	Thr	Leu	Phe	Pro	Phe	Glu	Ala	Glu	Ala	Tyr	Arg	Asn	Ile	Glu	Pro	
				140					145					150		
GTC	TAT	TTA	AAT	ATG	CTG	CTG	GAA	AAA	CTC	TGT	GTC	ATG	TTT	GAA	GAC	833
Val	Tyr	Leu	Asn	Met	Leu	Leu	Glu	Lys	Leu	Cys	Val	Met	Phe	Glu	Asp	
			155					160					165			
GGT	GTG	CTC	ATG	CGG	CTT	AAG	TCT	GAT	TTG	CTA	AAA	GCA	GCT	TTG	TGC	881
Gly	Val	Leu	Met	Arg	Leu	Lys	Ser	Asp	Leu	Leu	Lys	Ala	Ala	Leu	Cys	
		170					175					180				
CAT	TTA	CTG	CAG	TAT	TTC	CTT	AAA	TTT	GTG	CCA	GCT	GGG	TAT	GAA	TCT	929
His	Leu	Leu	Gln	Tyr	Phe	Leu	Lys	Phe	Val	Pro	Ala	Gly	Tyr	Glu	Ser	
	185					190					195					
GCT	TTA	CAA	GTC	AGG	AAG	GTC	TAT	GTG	AGA	AAT	ATT	TGT	AAA	GCT	CTT	977
Ala	Leu	Gln	Val	Arg	Lys	Val	Tyr	Val	Arg	Asn	Ile	Cys	Lys	Ala	Leu	
200					205					210					215	
TTG	GAT	GTG	CTT	GGA	ATT	GAG	GTA	GAT	GCA	GAG	TAC	TTG	TTG	GGC	CCA	1025
Leu	Asp	Val	Leu	Gly	Ile	Glu	Val	Asp	Ala	Glu	Tyr	Leu	Leu	Gly	Pro	
				220					225					230		
CTT	TAT	GCA	GCT	TTG	AAA	ATG	GAA	AGT	ATG	GAA	ATC	ATT	GAG	GAG	ATT	1073
Leu	Tyr	Ala	Ala	Leu	Lys	Met	Glu	Ser	Met	Glu	Ile	Ile	Glu	Glu	Ile	
			235					240					245			
CAA	TGC	CAA	ACT	CAA	CAG	GAA	AAC	CTC	AGC	AGT	AAT	AGT	GAT	GGA	ATA	1121
Gln	Cys	Gln	Thr	Gln	Gln	Glu	Asn	Leu	Ser	Ser	Asn	Ser	Asp	Gly	Ile	
		250					255					260				
TCA	CCC	AAA	AGG	CGT	CGT	CTC	AGC	TCG	TCT	CTA	AAC	CCT	TCT	AAA	AGA	1169
Ser	Pro	Lys	Arg	Arg	Arg	Leu	Ser	Ser	Ser	Leu	Asn	Pro	Ser	Lys	Arg	
	265					270					275					
GCA	CCA	AAA	CAG	ACT	GAG	GAA	ATT	AAA	CAT	GTG	GAC	ATG	AAC	CAA	AAG	1217
Ala	Pro	Lys	Gln	Thr	Glu	Glu	Ile	Lys	His	Val	Asp	Met	Asn	Gln	Lys	
280					285					290					295	
AGC	ATA	TTA	TGG	AGT	GCA	CTG	AAA	CAG	AAA	GCT	GAA	TCC	CTT	CAG	ATT	1265
Ser	Ile	Leu	Trp	Ser	Ala	Leu	Lys	Gln	Lys	Ala	Glu	Ser	Leu	Gln	Ile	
				300					305					310		

-150-

TCC	CTT	GAA	TAC	AGT	GGC	CTA	AAG	AAT	CCT	GTT	ATT	GAG	ATG	TTA	GAA	1313
Ser	Leu	Glu	Tyr	Ser	Gly	Leu	Lys	Asn	Pro	Val	Ile	Glu	Met	Leu	Glu	
		315						320					325			
GGA	ATT	GCT	GTT	GTC	TTA	CAA	CTG	ACT	GCT	CTG	TGT	ACT	GTT	CAT	TGT	1361
Gly	Ile	Ala	Val	Val	Leu	Gln	Leu	Thr	Ala	Leu	Cys	Thr	Val	His	Cys	
		330					335					340				
TCT	CAT	CAA	AAC	ATG	AAC	TGC	CGT	ACT	TTC	AAG	GAC	TGT	CAA	CAT	AAA	1409
Ser	His	Gln	Asn	Met	Asn	Cys	Arg	Thr	Phe	Lys	Asp	Cys	Gln	His	Lys	
	345					350					355					
TCC	AAG	AAG	AAA	CCT	TCT	GTA	GTG	ATA	ACT	TGG	ATG	TCA	TTG	GAT	TTT	1457
Ser	Lys	Lys	Lys	Pro	Ser	Val	Val	Ile	Thr	Trp	Met	Ser	Leu	Asp	Phe	
360					365					370					375	
TAC	ACA	AAA	GTG	CTT	AAG	AGC	TGT	AGA	AGT	TTG	TTA	GAA	TCT	GTT	CAG	1505
Tyr	Thr	Lys	Val	Leu	Lys	Ser	Cys	Arg	Ser	Leu	Leu	Glu	Ser	Val	Gln	
			380						385					390		
AAA	CTG	GAC	CTG	GAG	GCA	ACC	ATT	GAT	AAG	GTG	GTG	AAA	ATT	TAT	GAT	1553
Lys	Leu	Asp	Leu	Glu	Ala	Thr	Ile	Asp	Lys	Val	Val	Lys	Ile	Tyr	Asp	
		395						400					405			
GCT	TTG	ATT	TAT	ATG	CAA	GTA	AAC	AGT	TCA	TTT	GAA	GAT	CAT	ATC	CTG	1601
Ala	Leu	Ile	Tyr	Met	Gln	Val	Asn	Ser	Ser	Phe	Glu	Asp	His	Ile	Leu	
		410					415					420				
GAA	GAT	TTA	TGT	GGA	ATG	CTC	TCA	CTT	CCA	TGG	ATT	TAT	TCC	CAT	TCT	1649
Glu	Asp	Leu	Cys	Gly	Met	Leu	Ser	Leu	Pro	Trp	Ile	Tyr	Ser	His	Ser	
	425					430					435					
GAT	GAT	GGC	TGT	TTA	AAG	TTG	ACC	ACA	TTT	GCC	GCT	AAT	CTT	CTA	ACA	1697
Asp	Asp	Gly	Cys	Leu	Lys	Leu	Thr	Thr	Phe	Ala	Ala	Asn	Leu	Leu	Thr	
440					445					450					455	
TTA	AGC	TGT	AGG	ATT	TCA	GAT	AGC	TAT	TCA	CCA	CAG	GCA	CAA	TCA	CGA	1745
Leu	Ser	Cys	Arg	Ile	Ser	Asp	Ser	Tyr	Ser	Pro	Gln	Ala	Gln	Ser	Arg	
			460					465						470		
TGT	GTG	TTT	CTT	CTG	ACT	CTG	TTT	CCA	AGA	AGA	ATA	TTC	CTT	GAG	TGG	1793
Cys	Val	Phe	Leu	Leu	Thr	Leu	Phe	Pro	Arg	Arg	Ile	Phe	Leu	Glu	Trp	
			475					480					485			
AGA	ACA	GCA	GTT	TAC	AAC	TGG	GCC	CTG	CAG	AGC	TCC	CAT	GAA	GTA	ATC	1841
Arg	Thr	Ala	Val	Tyr	Asn	Trp	Ala	Leu	Gln	Ser	Ser	His	Glu	Val	Ile	
		490					495					500				
CGG	GCT	AGT	TGT	GTT	AGT	GGA	TTT	TTT	ATC	TTA	TTG	CAG	CAG	CAG	AAT	1889
Arg	Ala	Ser	Cys	Val	Ser	Gly	Phe	Phe	Ile	Leu	Leu	Gln	Gln	Gln	Asn	
	505					510						515				
TCT	TGT	AAC	AGA	GTT	CCC	AAG	ATT	CTT	ATA	GAT	AAA	GTC	AAA	GAT	GAT	1937
Ser	Cys	Asn	Arg	Val	Pro	Lys	Ile	Leu	Ile	Asp	Lys	Val	Lys	Asp	Asp	
520					525					530					535	
TCT	GAC	ATT	GTC	AAG	AAA	GAA	TTT	GCT	TCT	ATA	CTT	GGT	CAA	CTT	GTC	1985
Ser	Asp	Ile	Val	Lys	Lys	Glu	Phe	Ala	Ser	Ile	Leu	Gly	Gln	Leu	Val	
			540					545						550		
TGT	ACT	CTT	CAC	GGC	ATG	TTT	TAT	CTG	ACA	AGT	TCT	TTA	ACA	GAA	CCT	2033
Cys	Thr	Leu	His	Gly	Met	Phe	Tyr	Leu	Thr	Ser	Ser	Leu	Thr	Glu	Pro	
			555					560					565			
TTC	TCT	GAA	CAC	GGA	CAT	GTG	GAC	CTC	TTC	TGT	AGG	AAC	TTG	AAA	GCC	2081
Phe	Ser	Glu	His	Gly	His	Val	Asp	Leu	Phe	Cys	Arg	Asn	Leu	Lys	Ala	
		570					575					580				

-151-

ACT	TCT	CAA	CAT	GAA	TGT	TCA	TCT	TCT	CAA	CTA	AAA	GCT	TCT	GTC	TGC	2129
Thr	Ser	Gln	His	Glu	Cys	Ser	Ser	Ser	Gln	Leu	Lys	Ala	Ser	Val	Cys	
585						590					595					
AAG	CCA	TTC	CTT	TTC	CTA	CTG	AAA	AAA	AAA	ATA	CCT	AGT	CCA	GTA	AAA	2177
Lys	Pro	Phe	Leu	Phe	Leu	Leu	Lys	Lys	Lys	Ile	Pro	Ser	Pro	Val	Lys	
600					605					610					615	
CTT	GCT	TTC	ATA	GAT	AAT	CTA	CAT	CAT	CTT	TGT	AAG	CAT	CTT	GAT	TTT	2225
Leu	Ala	Phe	Ile	Asp	Asn	Leu	His	His	Leu	Cys	Lys	His	Leu	Asp	Phe	
				620					625					630		
AGA	GAA	GAT	GAA	ACA	GAT	GTA	AAA	GCA	GTT	CTT	GGA	ACT	TTA	TTA	AAT	2273
Arg	Glu	Asp	Glu	Thr	Asp	Val	Lys	Ala	Val	Leu	Gly	Thr	Leu	Leu	Asn	
			635					640					645			
TTA	ATG	GAA	GAT	CCA	GAC	AAA	GAT	GTT	AGA	GTG	GCT	TTT	AGT	GGA	AAT	2321
Leu	Met	Glu	Asp	Pro	Asp	Lys	Asp	Val	Arg	Val	Ala	Phe	Ser	Gly	Asn	
		650					655					660				
ATC	AAG	CAC	ATA	TTG	GAA	TCC	TTG	GAC	TCT	GAA	GAT	GGA	TTT	ATA	AAG	2369
Ile	Lys	His	Ile	Leu	Glu	Ser	Leu	Asp	Ser	Glu	Asp	Gly	Phe	Ile	Lys	
	665					670					675					
GAG	CTT	TTT	GTC	TTA	AGA	ATG	AAG	GAA	GCA	TAT	ACA	CAT	GCC	CAA	ATA	2417
Glu	Leu	Phe	Val	Leu	Arg	Met	Lys	Glu	Ala	Tyr	Thr	His	Ala	Gln	Ile	
680					685					690					695	
TCA	AGA	AAT	AAT	GAG	CTG	AAG	GAT	ACC	TTG	ATT	CTT	ACA	ACA	GGG	GAT	2465
Ser	Arg	Asn	Asn	Glu	Leu	Lys	Asp	Thr	Leu	Ile	Leu	Thr	Thr	Gly	Asp	
				700					705					710		
ATT	GGA	AGG	GCC	GCA	AAA	GGA	GAT	TTG	GTA	CCA	TTT	GCA	CTC	TTA	CAC	2513
Ile	Gly	Arg	Ala	Ala	Lys	Gly	Asp	Leu	Val	Pro	Phe	Ala	Leu	Leu	His	
			715					720					725			
TTA	TTG	CAT	TGT	TTG	TTA	TCC	AAG	TCA	GCA	TCT	GTC	TCT	GGA	GCA	GCA	2561
Leu	Leu	His	Cys	Leu	Leu	Ser	Lys	Ser	Ala	Ser	Val	Ser	Gly	Ala	Ala	
		730					735					740				
TAC	ACA	GAA	ATT	AGA	GCT	CTG	GTT	GCA	GCT	AAA	AGT	GTT	AAA	CTG	CAA	2609
Tyr	Thr	Glu	Ile	Arg	Ala	Leu	Val	Ala	Ala	Lys	Ser	Val	Lys	Leu	Gln	
	745					750					755					
AGT	TTT	TTC	AGC	CAG	TAT	AAG	AAA	CCC	ATC	TGT	CAG	TTT	TTG	GTA	GAA	2657
Ser	Phe	Phe	Ser	Gln	Tyr	Lys	Lys	Pro	Ile	Cys	Gln	Phe	Leu	Val	Glu	
760					765					770					775	
TCC	CTT	CAC	TCT	AGT	CAG	ATG	ACA	GCA	CTT	CCG	AAT	ACT	CCA	TGC	CAG	2705
Ser	Leu	His	Ser	Ser	Gln	Met	Thr	Ala	Leu	Pro	Asn	Thr	Pro	Cys	Gln	
				780					785					790		
AAT	GCT	GAC	GTG	CGA	AAA	CAA	GAT	GTG	GCT	CAC	CAG	AGA	GAA	ATG	GCT	2753
Asn	Ala	Asp	Val	Arg	Lys	Gln	Asp	Val	Ala	His	Gln	Arg	Glu	Met	Ala	
			795					800					805			
TTA	AAT	ACG	TTG	TCT	GAA	ATT	GCC	AAC	GTT	TTC	GAC	TTT	CCT	GAT	CTT	2801
Leu	Asn	Thr	Leu	Ser	Glu	Ile	Ala	Asn	Val	Phe	Asp	Phe	Pro	Asp	Leu	
		810					815					820				
AAT	CGT	TTT	CTT	ACT	AGG	ACA	TTA	CAA	GTT	CTA	CTA	CCT	GAT	CTT	GCT	2849
Asn	Arg	Phe	Leu	Thr	Arg	Thr	Leu	Gln	Val	Leu	Leu	Pro	Asp	Leu	Ala	
	825					830					835					
GCC	AAA	GCA	AGC	CCT	GCA	GCT	TCT	GCT	CTC	ATT	CGA	ACT	TTA	GGA	AAA	2897
Ala	Lys	Ala	Ser	Pro	Ala	Ala	Ser	Ala	Leu	Ile	Arg	Thr	Leu	Gly	Lys	
840					845				850						855	

-152-

CAA	TTA	AAT	GTC	AAT	CGT	AGA	GAG	ATT	TTA	ATA	AAC	AAC	TTC	AAA	TAT	2945
Gln	Leu	Asn	Val	Asn	Arg	Arg	Glu	Ile	Leu	Ile	Asn	Asn	Phe	Lys	Tyr	
				860					865					870		
ATT	TTT	TCT	CAT	TTG	GTC	TGT	TCT	TGT	TCC	AAA	GAT	GAA	TTA	GAA	CGT	2993
Ile	Phe	Ser	His	Leu	Val	Cys	Ser	Cys	Ser	Lys	Asp	Glu	Leu	Glu	Arg	
			875					880					885			
GCC	CTT	CAT	TAT	CTG	AAG	AAT	GAA	ACA	GAA	ATT	GAA	CTG	GGG	AGC	CTG	3041
Ala	Leu	His	Tyr	Leu	Lys	Asn	Glu	Thr	Glu	Ile	Glu	Leu	Gly	Ser	Leu	
			890				895					900				
TTG	AGA	CAA	GAT	TTC	CAA	GGA	TTG	CAT	AAT	GAA	TTA	TTG	CTG	CGT	ATT	3089
Leu	Arg	Gln	Asp	Phe	Gln	Gly	Leu	His	Asn	Glu	Leu	Leu	Leu	Arg	Ile	
	905					910					915					
GGA	GAA	CAC	TAT	CAA	CAG	GTT	TTT	AAT	GGT	TTG	TCA	ATA	CTT	GCC	TCA	3137
Gly	Glu	His	Tyr	Gln	Gln	Val	Phe	Asn	Gly	Leu	Ser	Ile	Leu	Ala	Ser	
920					925					930					935	
TTT	GCA	TCC	AGT	GAT	GAT	CCA	TAT	CAG	GGC	CCG	AGA	GAT	ATC	ATA	TCA	3185
Phe	Ala	Ser	Ser	Asp	Asp	Pro	Tyr	Gln	Gly	Pro	Arg	Asp	Ile	Ile	Ser	
				940					945					950		
CCT	GAA	CTG	ATG	GCT	GAT	TAT	TTA	CAA	CCC	AAA	TTG	TTG	GGC	ATT	TTG	3233
Pro	Glu	Leu	Met	Ala	Asp	Tyr	Leu	Gln	Pro	Lys	Leu	Leu	Gly	Ile	Leu	
			955					960					965			
GCT	TTT	TTT	AAC	ATG	CAG	TTA	CTG	AGC	TCT	AGT	GTT	GGC	ATT	GAA	GAT	3281
Ala	Phe	Phe	Asn	Met	Gln	Leu	Leu	Ser	Ser	Ser	Val	Gly	Ile	Glu	Asp	
			970				975					980				
AAG	AAA	ATG	GCC	TTG	AAC	AGT	TTG	ATG	TCT	TTG	ATG	AAG	TTA	ATG	GGA	3329
Lys	Lys	Met	Ala	Leu	Asn	Ser	Leu	Met	Ser	Leu	Met	Lys	Leu	Met	Gly	
	985					990					995					
CCC	AAA	CAT	GTC	AGT	TCT	GTG	AGG	GTG	AAG	ATG	ATG	ACC	ACA	CTG	AGA	3377
Pro	Lys	His	Val	Ser	Ser	Val	Arg	Val	Lys	Met	Met	Thr	Thr	Leu	Arg	
1000					1005					1010					1015	
ACT	GGC	CTT	CGA	TTC	AAG	GAT	GAT	TTT	CCT	GAA	TTG	TGT	TGC	AGA	GCT	3425
Thr	Gly	Leu	Arg	Phe	Lys	Asp	Asp	Phe	Pro	Glu	Leu	Cys	Cys	Arg	Ala	
				1020					1025					1030		
TGG	GAC	TGC	TTT	GTT	CGC	TGC	CTG	GAT	CAT	GCT	TGT	CTG	GGC	TCC	CTT	3473
Trp	Asp	Cys	Phe	Val	Arg	Cys	Leu	Asp	His	Ala	Cys	Leu	Gly	Ser	Leu	
			1035					1040					1045			
CTC	AGT	CAT	GTA	ATA	GTA	GCT	TTG	TTA	CCT	CTT	ATA	CAC	ATC	CAG	CCT	3521
Leu	Ser	His	Val	Ile	Val	Ala	Leu	Leu	Pro	Leu	Ile	His	Ile	Gln	Pro	
			1050				1055					1060				
AAA	GAA	ACT	GCA	GCT	ATC	TTC	CAC	TAC	CTC	ATA	ATT	GAA	AAC	AGG	GAT	3569
Lys	Glu	Thr	Ala	Ala	Ile	Phe	His	Tyr	Leu	Ile	Ile	Glu	Asn	Arg	Asp	
	1065					1070					1075					
GCT	GTG	CAA	GAT	TTT	CTT	CAT	GAA	ATA	TAT	TTT	TTA	CCT	GAT	CAT	CCA	3617
Ala	Val	Gln	Asp	Phe	Leu	His	Glu	Ile	Tyr	Phe	Leu	Pro	Asp	His	Pro	
1080						1085				1090					1095	
GAA	TTA	AAA	AAG	ATA	AAA	GCC	GTT	CTC	CAG	GAA	TAC	AGA	AAG	GAG	ACC	3665
Glu	Leu	Lys	Lys	Ile	Lys	Ala	Val	Leu	Gln	Glu	Tyr	Arg	Lys	Glu	Thr	
				1100					1105					1110		
TCT	GAG	AGC	ACT	GAT	CTT	CAG	ACA	ACT	CTT	CAG	CTC	TCT	ATG	AAG	GCC	3713
Ser	Glu	Ser	Thr	Asp	Leu	Gln	Thr	Thr	Leu	Gln	Leu	Ser	Met	Lys	Ala	
			1115					1120					1125			

-153-

ATT CAA CAT GAA AAT GTC GAT GTT CGT ATT CAT GCT CTT ACA AGC TTG Ile Gln His Glu Asn Val Asp Val Arg Ile His Ala Leu Thr Ser Leu 1130 1135 1140	3761
AAG GAA ACC TTG TAT AAA AAT CAG GAA AAA CTG ATA AAG TAT GCA ACA Lys Glu Thr Leu Tyr Lys Asn Gln Glu Lys Leu Ile Lys Tyr Ala Thr 1145 1150 1155	3809
GAC AGT GAA ACA GTA GAA CCT ATT ATC TCA CAG TTG GTG ACA GTG CTT Asp Ser Glu Thr Val Glu Pro Ile Ile Ser Gln Leu Val Thr Val Leu 1160 1165 1170 1175	3857
TTG AAA GGT TGC CAA GAT GCA AAC TCT CAA GCT CGG TTG CTC TGT GGG Leu Lys Gly Cys Gln Asp Ala Asn Ser Gln Ala Arg Leu Leu Cys Gly 1180 1185 1190	3905
GAA TGT TTA GGG GAA TTG GGG GCG ATA GAT CCA GGT CGA TTA GAT TTC Glu Cys Leu Glu Leu Gly Ala Ile Asp Pro Gly Arg Leu Asp Phe 1195 1200 1205	3953
TCA ACA ACT GAA ACT CAA GGA AAA GAT TTT ACA TTT GTG ACT GGA GTA Ser Thr Thr Glu Thr Gln Gly Lys Asp Phe Thr Phe Val Thr Gly Val 1210 1215 1220	4001
GAA GAT TCA AGC TTT GCC TAT GGA TTA TTG ATG GAG CTA ACA AGA GCT Glu Asp Ser Ser Phe Ala Tyr Gly Leu Leu Met Glu Leu Thr Arg Ala 1225 1230 1235	4049
TAC CTT GCG TAT GCT GAT AAT AGC CGA GCT CAA GAT TCA GCT GCC TAT Tyr Leu Ala Tyr Ala Asp Asn Ser Arg Ala Gln Asp Ser Ala Ala Tyr 1240 1245 1250 1255	4097
GCC ATT CAG GAG TTG CTT TCT ATT TAT GAC TGT AGA GAG ATG GAG ACC Ala Ile Gln Glu Leu Leu Ser Ile Tyr Asp Cys Arg Glu Met Glu Thr 1260 1265 1270	4145
AAC GGC CCA GGT CAC CAA TTG TGG AGG AGA TTT CCT GAG CAT GTT CGG Asn Gly Pro Gly His Gln Leu Trp Arg Arg Phe Pro Glu His Val Arg 1275 1280 1285	4193
GAA ATA CTA GAA CCT CAT CTA AAT ACC AGA TAC AAG AGT TCT CAG AAG Glu Ile Leu Glu Pro His Leu Asn Thr Arg Tyr Lys Ser Ser Gln Lys 1290 1295 1300	4241
TCA ACC GAT TGG TCT GGA GTA AAG AAG CCA ATT TAC TTA AGT AAA TTG Ser Thr Asp Trp Ser Gly Val Lys Lys Pro Ile Tyr Leu Ser Lys Leu 1305 1310 1315	4289
GGT AGT AAC TTT GCA GAA TGG TCA GCA TCT TGG GCA GGT TAT CTT ATT Gly Ser Asn Phe Ala Glu Trp Ser Ala Ser Trp Ala Gly Tyr Leu Ile 1320 1325 1330 1335	4337
ACA AAG GTT CGA CAT GAT CTT GCC AGT AAA ATT TTC ACC TGC TGT AGC Thr Lys Val Arg His Asp Leu Ala Ser Lys Ile Phe Thr Cys Cys Ser 1340 1345 1350	4385
ATT ATG ATG AAG CAT GAT TTC AAA GTG ACC ATC TAT CTT CTT CCA CAT Ile Met Met Lys His Asp Phe Lys Val Thr Ile Tyr Leu Leu Pro His 1355 1360 1365	4433
ATT CTG GTG TAT GTC TTA CTG GGT TGT AAT CAA GAA GAT CAG CAG GAG Ile Leu Val Tyr Val Leu Leu Gly Cys Asn Gln Glu Asp Gln Gln Glu 1370 1375 1380	4481
GTT TAT GCA GAA ATT ATG GCA GTT CTA AAG CAT GAC GAT CAG CAT ACC Val Tyr Ala Glu Ile Met Ala Val Leu Lys His Asp Asp Gln His Thr 1385 1390 1395	4529

-154-

ATA AAT ACC CAA GAC ATT GCA TCT GAT CTG TGT CAA CTC AGT ACA CAG Ile Asn Thr Gln Asp Ile Ala Ser Asp Leu Cys Gln Leu Ser Thr Gln 1400 1405 1410 1415	4577
ACT GTG TTC TCC ATG CTT GAC CAT CTC ACA CAG TGG GCA AGG CAC AAA Thr Val Phe Ser Met Leu Asp His Leu Thr Gln Trp Ala Arg His Lys 1420 1425 1430	4625
TTT CAG GCA CTG AAA GCT GAG AAA TGT CCA CAC AGC AAA TCA AAC AGA Phe Gln Ala Leu Lys Ala Glu Lys Cys Pro His Ser Lys Ser Asn Arg 1435 1440 1445	4673
AAT AAG GTA GAC TCA ATG GTA TCT ACT GTG GAT TAT GAA GAC TAT CAG Asn Lys Val Asp Ser Met Val Ser Thr Val Asp Tyr Glu Asp Tyr Gln 1450 1455 1460	4721
AGT GTA ACC CGT TTT CTA GAC CTC ATA CCC CAG GAT ACT CTG GCA GTA Ser Val Thr Arg Phe Leu Asp Leu Ile Pro Gln Asp Thr Leu Ala Val 1465 1470 1475	4769
GCT TCC TTT CGC TCC AAA GCA TAC ACA CGA GCT GTA ATG CAC TTT GAA Ala Ser Phe Arg Ser Lys Ala Tyr Thr Arg Ala Val Met His Phe Glu 1480 1485 1490 1495	4817
TCA TTT ATT ACA GAA AAG AAG CAA AAT ATT CAG GAA CAT CTT GGA TTT Ser Phe Ile Thr Glu Lys Lys Gln Asn Ile Gln Glu His Leu Gly Phe 1500 1505 1510	4865
TTA CAG AAA TTG TAT GCT GCT ATG CAT GAA CCT GAT GGA GTG GCC GGA Leu Gln Lys Leu Tyr Ala Ala Met His Glu Pro Asp Gly Val Ala Gly 1515 1520 1525	4913
GTC AGT GCA ATT AGA AAG GCA GAA CCA TCT CTA AAA GAA CAG ATC CTT Val Ser Ala Ile Arg Lys Ala Glu Pro Ser Leu Lys Glu Gln Ile Leu 1530 1535 1540	4961
GAA CAT GAA AGC CTT GGC TTG CTG AGG GAT GCC ACT GCT TGT TAT GAC Glu His Glu Ser Leu Gly Leu Leu Arg Asp Ala Thr Ala Cys Tyr Asp 1545 1550 1555	5009
AGG GCT ATT CAG CTA GAA CCA GAC CAG ATC ATT CAT TAC CAT GGT GTA Arg Ala Ile Gln Leu Glu Pro Asp Gln Ile Ile His Tyr His Gly Val 1560 1565 1570 1575	5057
GTA AAG TCC ATG TTA GGT CTT GGT CAG CTG TCT ACT GTT ATC ACT CAG Val Lys Ser Met Leu Gly Leu Gly Gln Leu Ser Thr Val Ile Thr Gln 1580 1585 1590	5105
GTG AAT GGA GTG CAT GCT AAC AGG TCC GAG TGG ACA GAT GAA TTA AAC Val Asn Gly Val His Ala Asn Arg Ser Glu Trp Thr Asp Glu Leu Asn 1595 1600 1605	5153
ACG TAC AGA GTG GAA GCA GCT TGG AAA TTG TCA CAG TGG GAT TTG GTG Thr Tyr Arg Val Glu Ala Ala Trp Lys Leu Ser Gln Trp Asp Leu Val 1610 1615 1620	5201
GAA AAC TAT TTG GCA GCA GAT GGA AAA TCT ACA ACA TGG AGT GTC AGA Glu Asn Tyr Leu Ala Ala Asp Gly Lys Ser Thr Thr Trp Ser Val Arg 1625 1630 1635	5249
CTG GGA CAG CTA TTA TTA TCA GCC AAA AAA AGA GAT ATC ACA GCT TTT Leu Gly Gln Leu Leu Leu Ser Ala Lys Lys Arg Asp Ile Thr Ala Phe 1640 1645 1650 1655	5297
TAT GAC TCA CTG AAA CTA GTG AGA GCA GAA CAA ATT GTA CCT CTT TCA Tyr Asp Ser Leu Lys Leu Val Arg Ala Glu Gln Ile Val Pro Leu Ser 1660 1665 1670	5345



-155-

GCT GCA AGC TTT GAA AGA GGC TCC TAC CAA CGA GGA TAT GAA TAT ATT	5393
Ala Ala Ser Phe Glu Arg Gly Ser Tyr Gln Arg Gly Tyr Glu Tyr Ile	
1675 1680 1685	
GTG AGA TTG CAC ATG TTA TGT GAG TTG GAG CAT AGC ATC AAA CCA CTT	5441
Val Arg Leu His Met Leu Cys Glu Leu Glu His Ser Ile Lys Pro Leu	
1690 1695 1700	
TTC CAG CAT TCT CCA GGT GAC AGT TCT CAA GAA GAT TCT CTA AAC TGG	5489
Phe Gln His Ser Pro Gly Asp Ser Ser Gln Glu Asp Ser Leu Asn Trp	
1705 1710 1715	
GTA GCT CGA CTA GAA ATG ACC CAG AAT TCC TAC AGA GCC AAG GAG CCT	5537
Val Ala Arg Leu Glu Met Thr Gln Asn Ser Tyr Arg Ala Lys Glu Pro	
1720 1725 1730 1735	
ATC CTG GCT CTC CGG AGG GCT TTA CTA AGC CTC AAC AAA AGA CCA GAT	5585
Ile Leu Ala Leu Arg Arg Ala Leu Leu Ser Leu Asn Lys Arg Pro Asp	
1740 1745 1750	
TAC AAT GAA ATG GTT GGA GAA TGC TGG CTG CAG AGT GCC AGG GTA GCT	5633
Tyr Asn Glu Met Val Gly Glu Cys Trp Leu Gln Ser Ala Arg Val Ala	
1755 1760 1765	
AGA AAG GCT GGT CAC CAC CAG ACA GCC TAC AAT GCT CTC CTT AAT GCA	5681
Arg Lys Ala Gly His His Gln Thr Ala Tyr Asn Ala Leu Leu Asn Ala	
1770 1775 1780	
GGG GAA TCA CGA CTC GCT GAA CTG TAC GTG GAA AGG GCA AAG TGG CTC	5729
Gly Glu Ser Arg Leu Ala Glu Leu Tyr Val Glu Arg Ala Lys Trp Leu	
1785 1790 1795	
TGG TCC AAG GGT GAT GTT CAC CAG GCA CTA ATT GTT CTT CAA AAA GGT	5777
Trp Ser Lys Gly Asp Val His Gln Ala Leu Ile Val Leu Gln Lys Gly	
1800 1805 1810 1815	
GTT GAA TTA TGT TTT CCT GAA AAT GAA ACC CCA CCT GAG GGT AAG AAC	5825
Val Glu Leu Cys Phe Pro Glu Asn Glu Thr Pro Pro Glu Gly Lys Asn	
1820 1825 1830	
ATG TTA ATC CAT GGT CGA GCT ATG CTA CTA GTG GGC CGA TTT ATG GAA	5873
Met Leu Ile His Gly Arg Ala Met Leu Leu Val Gly Arg Phe Met Glu	
1835 1840 1845	
GAA ACA GCT AAC TTT GAA AGC AAT GCA ATT ATG AAA AAA TAT AAG GAT	5921
Glu Thr Ala Asn Phe Glu Ser Asn Ala Ile Met Lys Lys Tyr Lys Asp	
1850 1855 1860	
GTG ACC GCG TGC CTG CCA GAA TGG GAG GAT GGG CAT TTT TAC CTT GCC	5969
Val Thr Ala Cys Leu Pro Glu Trp Glu Asp Gly His Phe Tyr Leu Ala	
1865 1870 1875	
AAG TAC TAT GAC AAA TTG ATG CCC ATG GTC ACA GAC AAC AAA ATG GAA	6017
Lys Tyr Tyr Asp Lys Leu Met Pro Met Val Thr Asp Asn Lys Met Glu	
1880 1885 1890 1895	
AAG CAA GGT GAT CTC ATC CGG TAT ATA GTT CTT CAT TTT GGC AGA TCT	6065
Lys Gln Gly Asp Leu Ile Arg Tyr Ile Val Leu His Phe Gly Arg Ser	
1900 1905 1910	
CTA CAA TAT GGA AAT CAG TTC ATA TAT CAG TCA ATG CCA CGA ATG TTA	6113
Leu Gln Tyr Gly Asn Gln Phe Ile Tyr Gln Ser Met Pro Arg Met Leu	
1915 1920 1925	
ACT CTA TGG CTT GAT TAT GGT ACA AAG GCA TAT GAA TGG GAA AAA GCT	6161
Thr Leu Trp Leu Asp Tyr Gly Thr Lys Ala Tyr Glu Trp Glu Lys Ala	
1930 1935 1940	

-156-

GGC Gly 1945	CGC Arg	TCC Ser	GAT Asp	CGT Arg	GTA Val	CAA Gln 1950	ATG Met	AGG Arg	AAT Asn	GAT Asp	TTG Leu 1955	GGT Gly	AAA Lys	ATA Ile	AAC Asn	6209
AAG Lys 1960	GTT Val	ATC Ile	ACA Thr	GAG Glu	CAT His 1965	ACA Thr	AAC Asn	TAT Tyr	TTA Leu	GCT Ala 1970	CCA Pro	TAT Tyr	CAA Gln	TTT Phe	TTG Leu 1975	6257
ACT Thr	GCT Ala	TTT Phe	TCA Ser	CAA Gln 1980	TTG Leu	ATC Ile	TCT Ser	CGA Arg	ATT Ile 1985	TGT Cys	CAT His	TCT Ser	CAC His	GAT Asp 1990	GAA Glu	6305
GTT Val	TTT Phe	GTT Val	GTC Val 1995	TTG Leu	ATG Met	GAA Glu	ATA Ile	ATA Ile 2000	GCC Ala	AAA Lys	GTA Val	TTT Phe	CTA Leu 2005	GCC Ala	TAT Tyr	6353
CCT Pro	CAA Gln 2010	CAA Gln	GCA Ala	ATG Met	TGG Trp	ATG Met	ATG Met 2015	ACA Thr	GCT Ala	GTG Val	TCA Ser	AAG Lys 2020	TCA Ser	TCT Ser	TAT Tyr	6401
CCC Pro 2025	ATG Met	CGT Arg	GTG Val	AAC Asn	AGA Arg	TGC Cys 2030	AAG Lys	GAA Glu	ATC Ile	CTC Leu	AAT Asn 2035	AAA Lys	GCT Ala	ATT Ile	CAT His	6449
ATG Met 2040	AAA Lys	AAA Lys	TCC Ser	TTA Leu	GAG Glu 2045	AAG Lys	TTT Phe	GTT Val	GGA Gly	GAT Asp 2050	GCA Ala	ACT Thr	CGC Arg	CTA Leu	ACA Thr 2055	6497
GAT Asp	AAG Lys	CTT Leu	CTA Leu	GAA Glu 2060	TTG Leu	TGC Cys	AAT Asn	AAA Lys	CCG Pro 2065	GTG Val	GAA Glu	ATT Ile	CTT Leu	GCT Ala 2070	TCT Ser	6545
CTT Leu	CAG Gln	AAA Lys 2075	CCA Pro	AAG Lys	AAG Lys	ATT Ile	TCT Ser	TTA Leu 2080	AAA Lys	GGC Gly	TCA Ser	GAT Asp	GGA Gly 2085	AAG Lys	TTC Phe	6593
TAC Tyr	ATC Ile	ATG Met 2090	ATG Met	TGT Cys	AAG Lys	CCA Pro	AAA Lys 2095	GAT Asp	GAC Asp	CTG Leu	AGA Arg	AAG Lys 2100	GAT Asp	TGT Cys	AGA Arg	6641
CTA Leu 2105	ATG Met	GAA Glu	TTC Phe	AAT Asn	TCC Ser	TTG Leu 2110	ATT Ile	AAT Asn	AAG Lys	TGC Cys 2115	TTA Arg	AGA Lys	AAA Lys	GAT Asp	GCA Ala	6689
GAG Glu 2120	TCT Ser	CGT Arg	AGA Arg	AGA Arg	GAA Glu 2125	CTT Leu	CAT His	ATT Ile	CGA Arg	ACA Thr 2130	TAT Tyr	GCA Ala	GTT Val	ATT Ile	CCA Pro 2135	6737
CTA Leu	AAT Asn	GAT Asp	GAA Glu	TGT Cys 2140	GGG Gly	ATT Ile	ATT Ile	GAA Glu	TGG Trp 2145	GTG Val	AAC Asn	AAC Asn	ACT Thr	GCT Ala 2150	GGT Gly	6785
TTG Leu	AGA Arg	CCT Pro	ATT Ile 2155	CTG Leu	ACC Thr	AAA Lys	CTA Leu 2160	TAT Tyr	AAA Lys	GAA Glu	AAG Lys	GGA Gly	GTG Val 2165	TAT Tyr	ATG Met	6833
ACA Thr	GGA Gly 2170	AAA Lys	GAA Glu	CTT Leu	CGC Arg	CAG Gln	TGT Cys 2175	ATG Met	CTA Leu	CCA Pro	AAG Lys	TCA Ser 2180	GCA Ala	GCT Ala	TTA Leu	6881
TCT Ser 2185	GAA Glu	AAA Lys	CTC Leu	AAA Lys	GTA Val	TTC Phe 2190	CGA Arg	GAA Glu	TTT Phe	CTC Leu 2195	CTG Leu	CCC Pro	AGG Arg	CAT His	CCT Pro	6929
CCT Pro 2200	ATT Ile	TTT Phe	CAT His	GAG Glu	TGG Trp 2205	TTT Phe	CTG Leu	AGA Arg	ACA Thr	TTC Phe 2210	CCT Pro	GAT Asp	CCT Pro	ACA Thr	TCA Ser 2215	6977

-157-

TGG TAC AGT AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA Trp Tyr Ser Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser 2220 2225 2230	7025
ATG GTT GGT TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT Met Val Gly Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile 2235 2240 2245	7073
CTC TTT GAT TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT Leu Phe Asp Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys 2250 2255 2260	7121
CTT TTC AAT AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT Leu Phe Asn Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe 2265 2270 2275	7169
CGC CTG ACT CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG Arg Leu Thr His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu 2280 2285 2290 2295	7217
GGT CTT TTT CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT Gly Leu Phe Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp 2300 2305 2310	7265
CAG CGA GAG CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT Gln Arg Glu Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro 2315 2320 2325	7313
CTT GTG GAA TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG Leu Val Glu Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu 2330 2335 2340	7361
AAT GAA ACT GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT Asn Glu Thr Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu 2345 2350 2355	7409
GAC ATT GAG CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG Asp Ile Glu Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val 2360 2365 2370 2375	7457
ACA GGA CTG CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA Thr Gly Leu Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln 2380 2385 2390	7505
GAA GCT ACT GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT Glu Ala Thr Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr 2395 2400 2405	7553
CCA TAT ATG TGAAATGAAA TTATGTAAAA GAATATGTTA ATAATCTAAA Pro Tyr Met 2410	7602
AGTAAAAAAAA AAAAAAAAAA AA	7624

## (2) INFORMATION FOR SEQ ID NO:31:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2410 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

-158-

Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser  
 1 5 10 15  
 Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu  
 20 25 30  
 Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr Leu Leu Met Val  
 35 40 45  
 Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu  
 50 55 60  
 Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile  
 65 70 75 80  
 Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly  
 85 90 95  
 Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu  
 100 105 110  
 Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu  
 115 120 125  
 Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu  
 130 135 140  
 Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys  
 145 150 155 160  
 Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp  
 165 170 175  
 Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe  
 180 185 190  
 Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val  
 195 200 205  
 Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly Ile Glu Val Asp  
 210 215 220  
 Ala Glu Tyr Leu Leu Gly Pro Leu Tyr Ala Ala Leu Lys Met Glu Ser  
 225 230 235 240  
 Met Glu Ile Ile Glu Glu Ile Gln Cys Gln Thr Gln Gln Glu Asn Leu  
 245 250 255  
 Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg Arg Leu Ser Ser  
 260 265 270  
 Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr Glu Glu Ile Lys  
 275 280 285  
 His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser Ala Leu Lys Gln  
 290 295 300  
 Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser Gly Leu Lys Asn  
 305 310 315 320  
 Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val Leu Gln Leu Thr  
 325 330 335  
 Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met Asn Cys Arg Thr  
 340 345 350

-159-

Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro Ser Val Val Ile  
 355 360 365  
 Thr Trp Met Ser Leu Asp Phe Tyr Thr Lys Val Leu Lys Ser Cys Arg  
 370 375 380  
 Ser Leu Leu Glu Ser Val Gln Lys Leu Asp Leu Glu Ala Thr Ile Asp  
 385 390 395 400  
 Lys Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met Gln Val Asn Ser  
 405 410 415  
 Ser Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly Met Leu Ser Leu  
 420 425 430  
 Pro Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu Lys Leu Thr Thr  
 435 440 445  
 Phe Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile Ser Asp Ser Tyr  
 450 455 460  
 Ser Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu Thr Leu Phe Pro  
 465 470 475 480  
 Arg Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr Asn Trp Ala Leu  
 485 490 495  
 Gln Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe  
 500 505 510  
 Ile Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu  
 515 520 525  
 Ile Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala  
 530 535 540  
 Ser Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu  
 545 550 555 560  
 Thr Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu  
 565 570 575  
 Phe Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser  
 580 585 590  
 Gln Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe Leu Leu Lys Lys  
 595 600 605  
 Lys Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp Asn Leu His His  
 610 615 620  
 Leu Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr Asp Val Lys Ala  
 625 630 635 640  
 Val Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro Asp Lys Asp Val  
 645 650 655  
 Arg Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu Glu Ser Leu Asp  
 660 665 670  
 Ser Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu Arg Met Lys Glu  
 675 680 685  
 Ala Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu Leu Lys Asp Thr  
 690 695 700

-160-

Leu Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala Lys Gly Asp Leu  
 705 710 715 720  
 Val Pro Phe Ala Leu Leu His Leu Leu His Cys Leu Leu Ser Lys Ser  
 725 730 735  
 Ala Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg Ala Leu Val Ala  
 740 745 750  
 Ala Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln Tyr Lys Lys Pro  
 755 760 765  
 Ile Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser Gln Met Thr Ala  
 770 775 780  
 Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val  
 785 790 795 800  
 Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn  
 805 810 815  
 Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln  
 820 825 830  
 Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala  
 835 840 845  
 Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile  
 850 855 860  
 Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys  
 865 870 875 880  
 Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu Lys Asn Glu Thr  
 885 890 895  
 Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe Gln Gly Leu His  
 900 905 910  
 Asn Glu Leu Leu Arg Ile Gly Glu His Tyr Gln Gln Val Phe Asn  
 915 920 925  
 Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp Asp Pro Tyr Gln  
 930 935 940  
 Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln  
 945 950 955 960  
 Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser  
 965 970 975  
 Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met  
 980 985 990  
 Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val  
 995 1000 1005  
 Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe  
 1010 1015 1020  
 Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp  
 1025 1030 1035 1040  
 His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu  
 1045 1050 1055

-161-

Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr  
 1060 1065 1070  
 Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile  
 1075 1080 1085  
 Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu  
 1090 1095 1100  
 Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr  
 1105 1110 1115 1120  
 Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg  
 1125 1130 1135  
 Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu  
 1140 1145 1150  
 Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile  
 1155 1160 1165  
 Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser  
 1170 1175 1180  
 Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile  
 1185 1190 1195 1200  
 Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp  
 1205 1210 1215  
 Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu  
 1220 1225 1230  
 Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg  
 1235 1240 1245  
 Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr  
 1250 1255 1260  
 Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg  
 1265 1270 1275 1280  
 Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr  
 1285 1290 1295  
 Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys  
 1300 1305 1310  
 Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala  
 1315 1320 1325  
 Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser  
 1330 1335 1340  
 Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val  
 1345 1350 1355 1360  
 Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys  
 1365 1370 1375  
 Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu  
 1380 1385 1390  
 Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp  
 1395 1400 1405

-162-

Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu  
 1410 1415 1420  
 Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys  
 1425 1430 1435 1440  
 Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr  
 1445 1450 1455  
 Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile  
 1460 1465 1470  
 Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr  
 1475 1480 1485  
 Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn  
 1490 1495 1500  
 Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His  
 1505 1510 1515 1520  
 Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg Lys Ala Glu Pro  
 1525 1530 1535  
 Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg  
 1540 1545 1550  
 Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln  
 1555 1560 1565  
 Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln  
 1570 1575 1580  
 Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser  
 1585 1590 1595 1600  
 Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys  
 1605 1610 1615  
 Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys  
 1620 1625 1630  
 Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys  
 1635 1640 1645  
 Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala  
 1650 1655 1660  
 Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr  
 1665 1670 1675 1680  
 Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu  
 1685 1690 1695  
 Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser  
 1700 1705 1710  
 Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn  
 1715 1720 1725  
 Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu  
 1730 1735 1740  
 Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp  
 1745 1750 1755 1760



-163-

Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala  
 1765 1770 1775  
 Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr  
 1780 1785 1790  
 Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala  
 1795 1800 1805  
 Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu  
 1810 1815 1820  
 Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu  
 1825 1830 1835 1840  
 Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala  
 1845 1850 1855  
 Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu  
 1860 1865 1870  
 Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met  
 1875 1880 1885  
 Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile  
 1890 1895 1900  
 Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr  
 1905 1910 1915 1920  
 Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys  
 1925 1930 1935  
 Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg  
 1940 1945 1950  
 Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr  
 1955 1960 1965  
 Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg  
 1970 1975 1980  
 Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Met Glu Ile Ile  
 1985 1990 1995 2000  
 Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr  
 2005 2010 2015  
 Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu  
 2020 2025 2030  
 Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val  
 2035 2040 2045  
 Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys  
 2050 2055 2060  
 Pro Val Glu Ile Leu Ala Ser Leu Gln Lys Pro Lys Lys Ile Ser Leu  
 2065 2070 2075 2080  
 Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met Met Cys Lys Pro Lys Asp  
 2085 2090 2095  
 Asp Leu Arg Lys Asp Cys Arg Leu Met Glu Phe Asn Ser Leu Ile Asn  
 2100 2105 2110

-164-

Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg Arg Arg Glu Leu His Ile  
 2115 2120 2125  
 Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp Glu Cys Gly Ile Ile Glu  
 2130 2135 2140  
 Trp Val Asn Asn Thr Ala Gly Leu Arg Pro Ile Leu Thr Lys Leu Tyr  
 2145 2150 2155 2160  
 Lys Glu Lys Gly Val Tyr Met Thr Gly Lys Glu Leu Arg Gln Cys Met  
 2165 2170 2175  
 Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys Leu Lys Val Phe Arg Glu  
 2180 2185 2190  
 Phe Leu Leu Pro Arg His Pro Pro Ile Phe His Glu Trp Phe Leu Arg  
 2195 2200 2205  
 Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser Ser Arg Ser Ala Tyr Cys  
 2210 2215 2220  
 Arg Ser Thr Ala Val Met Ser Met Val Gly Tyr Ile Leu Gly Leu Gly  
 2225 2230 2235 2240  
 Asp Arg His Gly Glu Asn Ile Leu Phe Asp Ser Leu Thr Gly Glu Cys  
 2245 2250 2255  
 Val His Val Asp Phe Asn Cys Leu Phe Asn Lys Gly Glu Thr Phe Glu  
 2260 2265 2270  
 Val Pro Glu Ile Val Pro Phe Arg Leu Thr His Asn Met Val Asn Gly  
 2275 2280 2285  
 Met Gly Pro Met Gly Thr Glu Gly Leu Phe Arg Arg Ala Cys Glu Val  
 2290 2295 2300  
 Thr Met Arg Leu Met Arg Asp Gln Arg Glu Pro Leu Met Ser Val Leu  
 2305 2310 2315 2320  
 Lys Thr Phe Leu His Asp Pro Leu Val Glu Trp Ser Lys Pro Val Lys  
 2325 2330 2335  
 Gly His Ser Lys Ala Pro Leu Asn Glu Thr Gly Glu Val Val Asn Glu  
 2340 2345 2350  
 Lys Ala Lys Thr His Val Leu Asp Ile Glu Gln Arg Leu Gln Gly Val  
 2355 2360 2365  
 Ile Lys Thr Arg Asn Arg Val Thr Gly Leu Pro Leu Ser Ile Glu Gly  
 2370 2375 2380  
 His Val His Tyr Leu Ile Gln Glu Ala Thr Asp Glu Asn Leu Leu Cys  
 2385 2390 2395 2400  
 Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2405 2410

## (2) INFORMATION FOR SEQ ID NO:32:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7502 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

-165-

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..7440

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

ATG GGT CAT GCT GTG GAA TGG CCA GTG GTC ATG AGC CGA TTT TTA AGT	48
Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser	
1 5 10 15	
CAA TTA GAT GAA CAC ATG GGA TAT TTA CAA TCA GCT CCT TTG CAG TTG	96
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu	
20 25 30	
ATG AGT ATG CAA AAT TTA GAA TTT ATT GAA GTC ACT TTA TTA ATG GTT	144
Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr Leu Leu Met Val	
35 40 45	
CTT ACT CGT ATT ATT GCA ATT GTG TTT TTT AGA AGG CAA GAA CTC TTA	192
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu	
50 55 60	
CTT TGG CAG ATA GGT TGT GTT CTG CTA GAG TAT GGT AGT CCA AAA ATT	240
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile	
65 70 75 80	
AAA TCC CTA GCA ATT AGC TTT TTA ACA GAA CTT TTT CAG CTT GGA GGA	288
Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly	
85 90 95	
CTA CCA GCA CAA CCA GCT AGC ACT TTT TTC AGC TCA TTT TTG GAA TTA	336
Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu	
100 105 110	
TTA AAA CAC CTT GTA GAA ATG GAT ACT GAC CAA TTG AAA CTC TAT GAA	384
Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu	
115 120 125	
GAG CCA TTA TCA AAG CTG ATA AAG ACA CTA TTT CCC TTT GAA GCA GAA	432
Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu	
130 135 140	
GCT TAT AGA AAT ATT GAA CCT GTC TAT TTA AAT ATG CTG CTG GAA AAA	480
Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys	
145 150 155 160	
CTC TGT GTC ATG TTT GAA GAC GGT GTG CTC ATG CGG CTT AAG TCT GAT	528
Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp	
165 170 175	
TTG CTA AAA GCA GCT TTG TGC CAT TTA CTG CAG TAT TTC CTT AAA TTT	576
Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe	
180 185 190	
GTG CCA GCT GGG TAT GAA TCT GCT TTA CAA GTC AGG AAG GTC TAT GTG	624
Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val	
195 200 205	
AGA AAT ATT TGT AAA GCT CTT TTG GAT GTG CTT GGA ATT GAG GTA GAT	672
Arg Asn Ile Cys Lys Ala Leu Asp Val Leu Gly Ile Glu Val Asp	
210 215 220	

-166-

GCA Ala 225	GAG Glu 225	TAC Tyr 225	TTG Leu 225	TTG Leu 225	GGC Gly 230	CCA Pro 230	CTT Leu 230	TAT Tyr 230	GCA Ala 235	GCT Ala 235	TTG Leu 235	AAA Lys 235	ATG Met 240	GAA Glu 240	AGT Ser 240	720
ATG Met 245	GAA Glu 245	ATC Ile 245	ATT Ile 245	GAG Glu 245	GAG Glu 245	ATT Ile 245	CAA Gln 250	TGC Cys 250	CAA Gln 250	ACT Thr 250	CAA Gln 250	CAG Gln 255	GAA Glu 255	AAC Asn 255	CTC Leu 255	768
AGC Ser 260	AGT Ser 260	AAT Asn 260	AGT Ser 260	GAT Asp 260	GGA Gly 265	ATA Ile 265	TCA Ser 265	CCC Pro 265	AAA Lys 265	AGG Arg 265	CGT Arg 265	CGT Arg 265	CTC Leu 270	AGC Ser 270	TCG Ser 270	816
TCT Ser 275	CTA Leu 275	AAC Asn 275	CCT Pro 275	TCT Ser 275	AAA Lys 280	AGA Arg 280	GCA Ala 280	CCA Pro 280	AAA Lys 280	CAG Gln 285	ACT Thr 285	GAG Glu 285	GAA Glu 285	ATT Ile 285	AAA Lys 285	864
CAT His 290	GTG Val 290	GAC Asp 290	ATG Met 290	AAC Asn 295	CAA Gln 295	AAG Lys 295	AGC Ser 295	ATA Ile 295	TTA Leu 300	TGG Trp 300	AGT Ser 300	GCA Ala 300	CTG Leu 300	AAA Lys 300	CAG Gln 300	912
AAA Lys 305	GCT Ala 305	GAA Glu 305	TCC Ser 305	CTT Leu 310	CAG Gln 310	ATT Ile 310	TCC Ser 310	CTT Leu 310	GAA Glu 315	TAC Tyr 315	AGT Ser 315	GGC Gly 315	CTA Leu 315	AAG Lys 315	AAT Asn 320	960
CCT Pro 325	GTT Val 325	ATT Ile 325	GAG Glu 325	ATG Met 325	TTA Leu 325	GAA Glu 325	GGA Gly 325	ATT Ile 330	GCT Ala 330	GTT Val 330	GTC Val 330	TTA Leu 330	CAA Gln 335	CTG Leu 335	ACT Thr 335	1008
GCT Ala 340	CTG Leu 340	TGT Cys 340	ACT Thr 340	GTT Val 340	CAT His 345	TGT Cys 345	TCT Ser 345	CAT His 345	CAA Gln 345	AAC Asn 345	ATG Met 345	AAC Asn 345	TGC Cys 350	CGT Arg 350	ACT Thr 350	1056
TTT Phe 355	AAG Lys 355	GAC Asp 355	TGT Cys 355	CAA Gln 360	CAT His 360	AAA Lys 360	TCC Ser 360	AAG Lys 360	AAG Lys 360	AAA Lys 360	CCT Pro 365	TCT Ser 365	GTA Val 365	GTG Val 365	ATA Ile 365	1104
ACT Thr 370	TGG Trp 370	ATG Met 370	TCA Ser 370	TTG Leu 375	GAT Asp 375	TTT Phe 375	TAC Tyr 375	ACA Thr 375	AAA Lys 380	GTG Val 380	CTT Leu 380	AAG Lys 380	AGC Ser 380	TGT Cys 380	AGA Arg 380	1152
AGT Ser 385	TTG Leu 385	TTA Leu 385	GAA Glu 385	TCT Ser 390	GTT Val 390	CAG Gln 390	AAA Lys 390	CTG Leu 395	GAC Asp 395	CTG Leu 395	GAG Glu 395	GCA Ala 395	ACC Thr 395	ATT Ile 395	GAT Asp 400	1200
AAG Lys 405	GTG Val 405	GTG Val 405	AAA Lys 405	ATT Ile 405	TAT Tyr 405	GAT Asp 410	GCT Ala 410	TTG Leu 410	ATT Ile 410	TAT Tyr 410	ATG Met 415	CAA Gln 415	GTA Val 415	AAC Asn 415	AGT Ser 415	1248
TCA Ser 420	TTT Phe 420	GAA Glu 420	GAT Asp 420	CAT His 425	ATC Ile 425	CTG Leu 425	GAA Glu 425	GAT Asp 425	TTA Leu 425	TGT Cys 425	GGA Gly 425	ATG Met 430	CTC Leu 430	TCA Ser 430	CTT Leu 430	1296
CCA Pro 435	TGG Trp 435	ATT Ile 435	TAT Tyr 435	TCC Ser 440	CAT His 440	TCT Ser 440	GAT Asp 440	GAT Asp 440	GGC Gly 445	TGT Cys 445	TTA Leu 445	AAG Lys 445	TTG Leu 445	ACC Thr 445	ACA Thr 445	1344
TTT Phe 450	GCC Ala 450	GCT Ala 450	AAT Asn 455	CTT Leu 455	CTA Leu 455	ACA Thr 455	TTA Leu 455	AGC Ser 460	TGT Cys 460	AGG Arg 460	ATT Ile 460	TCA Ser 460	GAT Asp 460	AGC Ser 460	TAT Tyr 460	1392
TCA Ser 465	CCA Pro 465	CAG Gln 465	GCA Ala 465	CAA Gln 470	TCA Ser 470	CGA Arg 470	TGT Cys 470	GTG Val 475	TTT Phe 475	CTT Leu 475	CTG Leu 475	ACT Thr 475	CTG Leu 475	TTT Phe 475	CCA Pro 480	1440
AGA Arg 485	AGA Arg 485	ATA Ile 485	TTC Phe 485	CTT Leu 485	GAG Glu 485	TGG Trp 485	AGA Arg 485	ACA Thr 490	GCA Ala 490	GTT Val 490	TAC Tyr 490	AAC Asn 490	TGG Trp 490	GCC Ala 490	CTG Leu 490	1488

-167-

CAG AGC TCC CAT GAA GTA ATC CGG GCT AGT TGT GTT AGT GGA TTT TTT	1536
Gln Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe	
500 505 510	
ATC TTA TTG CAG CAG CAG AAT TCT TGT AAC AGA GTT CCC AAG ATT CTT	1584
Ile Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu	
515 520 525	
ATA GAT AAA GTC AAA GAT GAT TCT GAC ATT GTC AAG AAA GAA TTT GCT	1632
Ile Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala	
530 535 540	
TCT ATA CTT GGT CAA CTT GTC TGT ACT CTT CAC GGC ATG TTT TAT CTG	1680
Ser Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu	
545 550 555 560	
ACA AGT TCT TTA ACA GAA CCT TTC TCT GAA CAC GGA CAT GTG GAC CTC	1728
Thr Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu	
565 570 575	
TTC TGT AGG AAC TTG AAA GCC ACT TCT CAA CAT GAA TGT TCA TCT TCT	1776
Phe Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser	
580 585 590	
CAA CTA AAA GCT TCT GTC TGC AAG CCA TTC CTT TTC CTA CTG AAA AAA	1824
Gln Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe Leu Leu Lys Lys	
595 600 605	
AAA ATA CCT AGT CCA GTA AAA CTT GCT TTC ATA GAT AAT CTA CAT CAT	1872
Lys Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp Asn Leu His His	
610 615 620	
CTT TGT AAG CAT CTT GAT TTT AGA GAA GAT GAA ACA GAT GTA AAA GCA	1920
Leu Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr Asp Val Lys Ala	
625 630 635 640	
GTT CTT GGA ACT TTA TTA AAT TTA ATG GAA GAT CCA GAC AAA GAT GTT	1968
Val Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro Asp Lys Asp Val	
645 650 655	
AGA GTG GCT TTT AGT GGA AAT ATC AAG CAC ATA TTG GAA TCC TTG GAC	2016
Arg Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu Glu Ser Leu Asp	
660 665 670	
TCT GAA GAT GGA TTT ATA AAG GAG CTT TTT GTC TTA AGA ATG AAG GAA	2064
Ser Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu Arg Met Lys Glu	
675 680 685	
GCA TAT ACA CAT GCC CAA ATA TCA AGA AAT AAT GAG CTG AAG GAT ACC	2112
Ala Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu Leu Lys Asp Thr	
690 695 700	
TTG ATT CTT ACA ACA GGG GAT ATT GGA AGG GCC GCA AAA GGA GAT TTG	2160
Leu Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala Lys Gly Asp Leu	
705 710 715 720	
GTA CCA TTT GCA CTC TTA CAC TTA TTG CAT TGT TTG TTA TCC AAG TCA	2208
Val Pro Phe Ala Leu Leu His Leu Leu His Cys Leu Leu Ser Lys Ser	
725 730 735	
GCA TCT GTC TCT GGA GCA GCA TAC ACA GAA ATT AGA GCT CTG GTT GCA	2256
Ala Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg Ala Leu Val Ala	
740 745 750	
GCT AAA AGT GTT AAA CTG CAA AGT TTT TTC AGC CAG TAT AAG AAA CCC	2304
Ala Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln Tyr Lys Lys Pro	
755 760 765	

-168-

ATC TGT CAG TTT TTG GTA GAA TCC CTT CAC TCT AGT CAG ATG ACA GCA Ile Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser Gln Met Thr Ala 770 775 780	2352
CTT CCG AAT ACT CCA TGC CAG AAT GCT GAC GTG CGA AAA CAA GAT GTG Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val 785 790 795 800	2400
GCT CAC CAG AGA GAA ATG GCT TTA AAT ACG TTG TCT GAA ATT GCC AAC Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn 805 810 815	2448
GTT TTC GAC TTT CCT GAT CTT AAT CGT TTT CTT ACT AGG ACA TTA CAA Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln 820 825 830	2496
GTT CTA CTA CCT GAT CTT GCT GCC AAA GCA AGC CCT GCA GCT TCT GCT Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala 835 840 845	2544
CTC ATT CGA ACT TTA GGA AAA CAA TTA AAT GTC AAT CGT AGA GAG ATT Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile 850 855 860	2592
TTA ATA AAC AAC TTC AAA TAT ATT TTT TCT CAT TTG GTC TGT TCT TGT Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys 865 870 875 880	2640
TCC AAA GAT GAA TTA GAA CGT GCC CTT CAT TAT CTG AAG AAT GAA ACA Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu Lys Asn Glu Thr 885 890 895	2688
GAA ATT GAA CTG GGG AGC CTG TTG AGA CAA GAT TTC CAA GGA TTG CAT Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe Gln Gly Leu His 900 905 910	2736
AAT GAA TTA TTG CTG CGT ATT GGA GAA CAC TAT CAA CAG GTT TTT AAT Asn Glu Leu Leu Arg Ile Gly Glu His Tyr Gln Gln Val Phe Asn 915 920 925	2784
GGT TTG TCA ATA CTT GCC TCA TTT GCA TCC AGT GAT GAT CCA TAT CAG Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp Asp Pro Tyr Gln 930 935 940	2832
GGC CCG AGA GAT ATC ATA TCA CCT GAA CTG ATG GCT GAT TAT TTA CAA Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln 945 950 955 960	2880
CCC AAA TTG TTG GGC ATT TTG GCT TTT TTT AAC ATG CAG TTA CTG AGC Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser 965 970 975	2928
TCT AGT GTT GGC ATT GAA GAT AAG AAA ATG GCC TTG AAC AGT TTG ATG Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met 980 985 990	2976
TCT TTG ATG AAG TTA ATG GGA CCC AAA CAT GTC AGT TCT GTG AGG GTG Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val 995 1000 1005	3024
AAG ATG ATG ACC ACA CTG AGA ACT GGC CTT CGA TTC AAG GAT GAT TTT Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe 1010 1015 1020	3072
CCT GAA TTG TGT TGC AGA GCT TGG GAC TGC TTT GTT CGC TGC CTG GAT Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp 1025 1030 1035 1040	3120

-169-

CAT GCT TGT CTG GGC TCC CTT CTC AGT CAT GTA ATA GTA GCT TTG TTA His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu 1045 1050 1055	3168
CCT CTT ATA CAC ATC CAG CCT AAA GAA ACT GCA GCT ATC TTC CAC TAC Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr 1060 1065 1070	3216
CTC ATA ATT GAA AAC AGG GAT GCT GTG CAA GAT TTT CTT CAT GAA ATA Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile 1075 1080 1085	3264
TAT TTT TTA CCT GAT CAT CCA GAA TTA AAA AAG ATA AAA GCC GTT CTC Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu 1090 1095 1100	3312
CAG GAA TAC AGA AAG GAG ACC TCT GAG AGC ACT GAT CTT CAG ACA ACT Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr 1105 1110 1115 1120	3360
CTT CAG CTC TCT ATG AAG GCC ATT CAA CAT GAA AAT GTC GAT GTT CGT Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg 1125 1130 1135	3408
ATT CAT GCT CTT ACA AGC TTG AAG GAA ACC TTG TAT AAA AAT CAG GAA Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu 1140 1145 1150	3456
AAA CTG ATA AAG TAT GCA ACA GAC AGT GAA ACA GTA GAA CCT ATT ATC Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile 1155 1160 1165	3504
TCA CAG TTG GTG ACA GTG CTT TTG AAA GGT TGC CAA GAT GCA AAC TCT Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser 1170 1175 1180	3552
CAA GCT CGG TTG CTC TGT GGG GAA TGT TTA GGG GAA TTG GGG GCG ATA Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile 1185 1190 1195 1200	3600
GAT CCA GGT CGA TTA GAT TTC TCA ACA ACT GAA ACT CAA GGA AAA GAT Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp 1205 1210 1215	3648
TTT ACA TTT GTG ACT GGA GTA GAA GAT TCA AGC TTT GCC TAT GGA TTA Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu 1220 1225 1230	3696
TTG ATG GAG CTA ACA AGA GCT TAC CTT GCG TAT GCT GAT AAT AGC CGA Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg 1235 1240 1245	3744
GCT CAA GAT TCA GCT GCC TAT GCC ATT CAG GAG TTG CTT TCT ATT TAT Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr 1250 1255 1260	3792
GAC TGT AGA GAG ATG GAG ACC AAC GGC CCA GGT CAC CAA TTG TGG AGG Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg 1265 1270 1275 1280	3840
AGA TTT CCT GAG CAT GTT CGG GAA ATA CTA GAA CCT CAT CTA AAT ACC Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr 1285 1290 1295	3888
AGA TAC AAG AGT TCT CAG AAG TCA ACC GAT TGG TCT GGA GTA AAG AAG Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys 1300 1305 1310	3936

-170-

CCA	ATT	TAC	TTA	AGT	AAA	TTG	GGT	AGT	AAC	TTT	GCA	GAA	TGG	TCA	GCA	3984
Pro	Ile	Tyr	Leu	Ser	Lys	Leu	Gly	Ser	Asn	Phe	Ala	Glu	Trp	Ser	Ala	
		1315					1320					1325				
TCT	TGG	GCA	GGT	TAT	CTT	ATT	ACA	AAG	GTT	CGA	CAT	GAT	CTT	GCC	AGT	4032
Ser	Trp	Ala	Gly	Tyr	Leu	Ile	Thr	Lys	Val	Arg	His	Asp	Leu	Ala	Ser	
	1330					1335				1340						
AAA	ATT	TTC	ACC	TGC	TGT	AGC	ATT	ATG	ATG	AAG	CAT	GAT	TTC	AAA	GTG	4080
Lys	Ile	Phe	Thr	Cys	Cys	Ser	Ile	Met	Met	Lys	His	Asp	Phe	Lys	Val	
1345					1350					1355					1360	
ACC	ATC	TAT	CTT	CTT	CCA	CAT	ATT	CTG	GTG	TAT	GTC	TTA	CTG	GGT	TGT	4128
Thr	Ile	Tyr	Leu	Leu	Pro	His	Ile	Leu	Val	Tyr	Val	Leu	Leu	Gly	Cys	
				1365					1370						1375	
AAT	CAA	GAA	GAT	CAG	CAG	GAG	GTT	TAT	GCA	GAA	ATT	ATG	GCA	GTT	CTA	4176
Asn	Gln	Glu	Asp	Gln	Gln	Glu	Val	Tyr	Ala	Glu	Ile	Met	Ala	Val	Leu	
			1380					1385					1390			
AAG	CAT	GAC	GAT	CAG	CAT	ACC	ATA	AAT	ACC	CAA	GAC	ATT	GCA	TCT	GAT	4224
Lys	His	Asp	Asp	Gln	His	Thr	Ile	Asn	Thr	Gln	Asp	Ile	Ala	Ser	Asp	
		1395					1400					1405				
CTG	TGT	CAA	CTC	AGT	ACA	CAG	ACT	GTG	TTC	TCC	ATG	CTT	GAC	CAT	CTC	4272
Leu	Cys	Gln	Leu	Ser	Thr	Gln	Thr	Val	Phe	Ser	Met	Leu	Asp	His	Leu	
	1410					1415					1420					
ACA	CAG	TGG	GCA	AGG	CAC	AAA	TTT	CAG	GCA	CTG	AAA	GCT	GAG	AAA	TGT	4320
Thr	Gln	Trp	Ala	Arg	His	Lys	Phe	Gln	Ala	Leu	Lys	Ala	Glu	Lys	Cys	
1425					1430					1435					1440	
CCA	CAC	AGC	AAA	TCA	AAC	AGA	AAT	AAG	GTA	GAC	TCA	ATG	GTA	TCT	ACT	4368
Pro	His	Ser	Lys	Ser	Asn	Arg	Asn	Lys	Val	Asp	Ser	Met	Val	Ser	Thr	
				1445					1450					1455		
GTG	GAT	TAT	GAA	GAC	TAT	CAG	AGT	GTA	ACC	CGT	TTT	CTA	GAC	CTC	ATA	4416
Val	Asp	Tyr	Glu	Asp	Tyr	Gln	Ser	Val	Thr	Arg	Phe	Leu	Asp	Leu	Ile	
			1460					1465					1470			
CCC	CAG	GAT	ACT	CTG	GCA	GTA	GCT	TCC	TTT	CGC	TCC	AAA	GCA	TAC	ACA	4464
Pro	Gln	Asp	Thr	Leu	Ala	Val	Ala	Ser	Phe	Arg	Ser	Lys	Ala	Tyr	Thr	
		1475					1480					1485				
CGA	GCT	GTA	ATG	CAC	TTT	GAA	TCA	TTT	ATT	ACA	GAA	AAG	AAG	CAA	AAT	4512
Arg	Ala	Val	Met	His	Phe	Glu	Ser	Phe	Ile	Thr	Glu	Lys	Lys	Gln	Asn	
	1490					1495					1500					
ATT	CAG	GAA	CAT	CTT	GGA	TTT	TTA	CAG	AAA	TTG	TAT	GCT	GCT	ATG	CAT	4560
Ile	Gln	Glu	His	Leu	Gly	Phe	Leu	Gln	Lys	Leu	Tyr	Ala	Ala	Met	His	
1505					1510					1515					1520	
GAA	CCT	GAT	GGA	GTG	GCC	GGA	GTC	AGT	GCA	ATT	AGA	AAG	GCA	GAA	CCA	4608
Glu	Pro	Asp	Gly	Val	Ala	Gly	Val	Ser	Ala	Ile	Arg	Lys	Ala	Glu	Pro	
				1525					1530					1535		
TCT	CTA	AAA	GAA	CAG	ATC	CTT	GAA	CAT	GAA	AGC	CTT	GGC	TTG	CTG	AGG	4656
Ser	Leu	Lys	Glu	Gln	Ile	Leu	Glu	His	Glu	Ser	Leu	Gly	Leu	Leu	Arg	
			1540					1545					1550			
GAT	GCC	ACT	GCT	TGT	TAT	GAC	AGG	GCT	ATT	CAG	CTA	GAA	CCA	GAC	CAG	4704
Asp	Ala	Thr	Ala	Cys	Tyr	Asp	Arg	Ala	Ile	Gln	Leu	Glu	Pro	Asp	Gln	
		1555					1560					1565				
ATC	ATT	CAT	TAC	CAT	GGT	GTA	GTA	AAG	TCC	ATG	TTA	GGT	CTT	GGT	CAG	4752
Ile	Ile	His	Tyr	His	Gly	Val	Val	Lys	Ser	Met	Leu	Gly	Leu	Gly	Gln	
	1570					1575					1580					



-171-

CTG TCT ACT GTT ATC ACT CAG GTG AAT GGA GTG CAT GCT AAC AGG TCC Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser 1585 1590 1595 1600	4800
GAG TGG ACA GAT GAA TTA AAC ACG TAC AGA GTG GAA GCA GCT TGG AAA Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys 1605 1610 1615	4848
TTG TCA CAG TGG GAT TTG GTG GAA AAC TAT TTG GCA GCA GAT GGA AAA Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys 1620 1625 1630	4896
TCT ACA ACA TGG AGT GTC AGA CTG GGA CAG CTA TTA TTA TCA GCC AAA Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Ser Ala Lys 1635 1640 1645	4944
AAA AGA GAT ATC ACA GCT TTT TAT GAC TCA CTG AAA CTA GTG AGA GCA Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala 1650 1655 1660	4992
GAA CAA ATT GTA CCT CTT TCA GCT GCA AGC TTT GAA AGA GGC TCC TAC Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr 1665 1670 1675 1680	5040
CAA CGA GGA TAT GAA TAT ATT GTG AGA TTG CAC ATG TTA TGT GAG TTG Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu 1685 1690 1695	5088
GAG CAT AGC ATC AAA CCA CTT TTC CAG CAT TCT CCA GGT GAC AGT TCT Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser 1700 1705 1710	5136
CAA GAA GAT TCT CTA AAC TGG GTA GCT CGA CTA GAA ATG ACC CAG AAT Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn 1715 1720 1725	5184
TCC TAC AGA GCC AAG GAG CCT ATC CTG GCT CTC CGG AGG GCT TTA CTA Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu 1730 1735 1740	5232
AGC CTC AAC AAA AGA CCA GAT TAC AAT GAA ATG GTT GGA GAA TGC TGG Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp 1745 1750 1755 1760	5280
CTG CAG AGT GCC AGG GTA GCT AGA AAG GCT GGT CAC CAC CAG ACA GCC Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala 1765 1770 1775	5328
TAC AAT GCT CTC CTT AAT GCA GGG GAA TCA CGA CTC GCT GAA CTG TAC Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr 1780 1785 1790	5376
GTG GAA AGG GCA AAG TGG CTC TGG TCC AAG GGT GAT GTT CAC CAG GCA Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala 1795 1800 1805	5424
CTA ATT GTT CTT CAA AAA GGT GTT GAA TTA TGT TTT CCT GAA AAT GAA Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu 1810 1815 1820	5472
ACC CCA CCT GAG GGT AAG AAC ATG TTA ATC CAT GGT CGA GCT ATG CTA Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu 1825 1830 1835 1840	5520
CTA GTG GGC CGA TTT ATG GAA GAA ACA GCT AAC TTT GAA AGC AAT GCA Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala 1845 1850 1855	5568

-172-

ATT ATG AAA AAA TAT AAG GAT GTG ACC GCG TGC CTG CCA GAA TGG GAG Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu 1860 1865 1870	5616
GAT GGG CAT TTT TAC CTT GCC AAG TAC TAT GAC AAA TTG ATG CCC ATG Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met 1875 1880 1885	5664
GTC ACA GAC AAC AAA ATG GAA AAG CAA GGT GAT CTC ATC CGG TAT ATA Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile 1890 1895 1900	5712
GTT CTT CAT TTT GGC AGA TCT CTA CAA TAT GGA AAT CAG TTC ATA TAT Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr 1905 1910 1915 1920	5760
CAG TCA ATG CCA CGA ATG TTA ACT CTA TGG CTT GAT TAT GGT ACA AAG Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys 1925 1930 1935	5808
GCA TAT GAA TGG GAA AAA GCT GGC CGC TCC GAT CGT GTA CAA ATG AGG Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg 1940 1945 1950	5856
AAT GAT TTG GGT AAA ATA AAC AAG GTT ATC ACA GAG CAT ACA AAC TAT Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr 1955 1960 1965	5904
TTA GCT CCA TAT CAA TTT TTG ACT GCT TTT TCA CAA TTG ATC TCT CGA Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg 1970 1975 1980	5952
ATT TGT CAT TCT CAC GAT GAA GTT TTT GTT GTC TTG ATG GAA ATA ATA Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Met Glu Ile Ile 1985 1990 1995 2000	6000
GCC AAA GTA TTT CTA GCC TAT CCT CAA CAA GCA ATG TGG ATG ATG ACA Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr 2005 2010 2015	6048
GCT GTG TCA AAG TCA TCT TAT CCC ATG CGT GTG AAC AGA TGC AAG GAA Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu 2020 2025 2030	6096
ATC CTC AAT AAA GCT ATT CAT ATG AAA AAA TCC TTA GAG AAG TTT GTT Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val 2035 2040 2045	6144
GGA GAT GCA ACT CGC CTA ACA GAT AAG CTT CTA GAA TTG TGC AAT AAA Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys 2050 2055 2060	6192
CCG GTT GAT GGA AGT AGT TCC ACA TTA AGC ATG AGC ACT CAT TTT AAA Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys 2065 2070 2075 2080	6240
ATG CTT AAA AAG CTG GTA GAA GAA GCA ACA TTT AGT GAA ATC CTC ATT Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile 2085 2090 2095	6288
CCT CTA CAA TCA GTC ATG ATA CCT ACA CTT CCA TCA ATT CTG GGT ACC Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr 2100 2105 2110	6336
CAT GCT AAC CAT GCT AGC CAT GAA CCA TTT CCT GGA CAT TGG GCC TAT His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr 2115 2120 2125	6384

-173-

ATT GCA GGG TTT GAT GAT ATG GTG GAA ATT CTT GCT TCT CTT CAG AAA Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys 2130 2135 2140	6432
CCA AAG AAG ATT TCT TTA AAA GGC TCA GAT GGA AAG TTC TAC ATC ATG Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met 2145 2150 2155 2160	6480
ATG TGT AAG CCA AAA GAT GAC CTG AGA AAG GAT TGT AGA CTA ATG GAA Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu 2165 2170 2175	6528
TTC AAT TCC TTG ATT AAT AAG TGC TTA AGA AAA GAT GCA GAG TCT CGT Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg 2180 2185 2190	6576
AGA AGA GAA CTT CAT ATT CGA ACA TAT GCA GTT ATT CCA CTA AAT GAT Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp 2195 2200 2205	6624
GAA TGT GGG ATT ATT GAA TGG GTG AAC AAC ACT GCT GGT TTG AGA CCT Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro 2210 2215 2220	6672
ATT CTG ACC AAA CTA TAT AAA GAA AAG GGA GTG TAT ATG ACA GGA AAA Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys 2225 2230 2235 2240	6720
GAA CTT CGC CAG TGT ATG CTA CCA AAG TCA GCA GCT TTA TCT GAA AAA Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys 2245 2250 2255	6768
CTC AAA GTA TTC CGA GAA TTT CTC CTG CCC AGG CAT CCT CCT ATT TTT Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro Ile Phe 2260 2265 2270	6816
CAT GAG TGG TTT CTG AGA ACA TTC CCT GAT CCT ACA TCA TGG TAC AGT His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser 2275 2280 2285	6864
AGT AGA TCA GCT TAC TGC CGT TCC ACT GCA GTA ATG TCA ATG GTT GGT Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly 2290 2295 2300	6912
TAT ATT CTG GGG CTT GGA GAC CGT CAT GGT GAA AAT ATT CTC TTT GAT Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp 2305 2310 2315 2320	6960
TCT TTG ACT GGT GAA TGC GTA CAT GTA GAT TTC AAT TGT CTT TTC AAT Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn 2325 2330 2335	7008
AAG GGA GAA ACC TTT GAA GTT CCA GAA ATT GTG CCA TTT CGC CTG ACT Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr 2340 2345 2350	7056
CAT AAT ATG GTT AAT GGA ATG GGT CCT ATG GGA ACA GAG GGT CTT TTT His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe 2355 2360 2365	7104
CGA AGA GCA TGT GAA GTT ACA ATG AGG CTG ATG CGT GAT CAG CGA GAG Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu 2370 2375 2380	7152
CCT TTA ATG AGT GTC TTA AAG ACT TTT CTA CAT GAT CCT CTT GTG GAA Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu 2385 2390 2395 2400	7200

-174-

TGG AGT AAA CCA GTG AAA GGG CAT TCC AAA GCG CCA CTG AAT GAA ACT	7248
Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr	
2405 2410 2415	
GGA GAA GTT GTC AAT GAA AAG GCC AAG ACC CAT GTT CTT GAC ATT GAG	7296
Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu	
2420 2425 2430	
CAG CGA CTA CAA GGT GTA ATC AAG ACT CGA AAT AGA GTG ACA GGA CTG	7344
Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu	
2435 2440 2445	
CCG TTA TCT ATT GAA GGA CAT GTG CAT TAC CTT ATA CAA GAA GCT ACT	7392
Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr	
2450 2455 2460	
GAT GAA AAC TTA CTA TGC CAG ATG TAT CTT GGT TGG ACT CCA TAT ATG	7440
Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met	
2465 2470 2475 2480	
TGAAATGAAA TTATGTAAAA GAATATGTTA ATAATCTAAA AGTAAAAAAAA AAAAAAAAAA	7500
AA	7502

## (2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2480 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

Met Gly His Ala Val Glu Trp Pro Val Val Met Ser Arg Phe Leu Ser	
1 5 10 15	
Gln Leu Asp Glu His Met Gly Tyr Leu Gln Ser Ala Pro Leu Gln Leu	
20 25 30	
Met Ser Met Gln Asn Leu Glu Phe Ile Glu Val Thr Leu Leu Met Val	
35 40 45	
Leu Thr Arg Ile Ile Ala Ile Val Phe Phe Arg Arg Gln Glu Leu Leu	
50 55 60	
Leu Trp Gln Ile Gly Cys Val Leu Leu Glu Tyr Gly Ser Pro Lys Ile	
65 70 75 80	
Lys Ser Leu Ala Ile Ser Phe Leu Thr Glu Leu Phe Gln Leu Gly Gly	
85 90 95	
Leu Pro Ala Gln Pro Ala Ser Thr Phe Phe Ser Ser Phe Leu Glu Leu	
100 105 110	
Leu Lys His Leu Val Glu Met Asp Thr Asp Gln Leu Lys Leu Tyr Glu	
115 120 125	
Glu Pro Leu Ser Lys Leu Ile Lys Thr Leu Phe Pro Phe Glu Ala Glu	
130 135 140	
Ala Tyr Arg Asn Ile Glu Pro Val Tyr Leu Asn Met Leu Leu Glu Lys	
145 150 155 160	

-175-

Leu Cys Val Met Phe Glu Asp Gly Val Leu Met Arg Leu Lys Ser Asp  
 165 170 175  
 Leu Leu Lys Ala Ala Leu Cys His Leu Leu Gln Tyr Phe Leu Lys Phe  
 180 185 190  
 Val Pro Ala Gly Tyr Glu Ser Ala Leu Gln Val Arg Lys Val Tyr Val  
 195 200 205  
 Arg Asn Ile Cys Lys Ala Leu Leu Asp Val Leu Gly Ile Glu Val Asp  
 210 215 220  
 Ala Glu Tyr Leu Leu Gly Pro Leu Tyr Ala Ala Leu Lys Met Glu Ser  
 225 230 235 240  
 Met Glu Ile Ile Glu Glu Ile Gln Cys Gln Thr Gln Gln Glu Asn Leu  
 245 250 255  
 Ser Ser Asn Ser Asp Gly Ile Ser Pro Lys Arg Arg Arg Leu Ser Ser  
 260 265 270  
 Ser Leu Asn Pro Ser Lys Arg Ala Pro Lys Gln Thr Glu Glu Ile Lys  
 275 280 285  
 His Val Asp Met Asn Gln Lys Ser Ile Leu Trp Ser Ala Leu Lys Gln  
 290 295 300  
 Lys Ala Glu Ser Leu Gln Ile Ser Leu Glu Tyr Ser Gly Leu Lys Asn  
 305 310 315 320  
 Pro Val Ile Glu Met Leu Glu Gly Ile Ala Val Val Leu Gln Leu Thr  
 325 330 335  
 Ala Leu Cys Thr Val His Cys Ser His Gln Asn Met Asn Cys Arg Thr  
 340 345 350  
 Phe Lys Asp Cys Gln His Lys Ser Lys Lys Lys Pro Ser Val Val Ile  
 355 360 365  
 Thr Trp Met Ser Leu Asp Phe Tyr Thr Lys Val Leu Lys Ser Cys Arg  
 370 375 380  
 Ser Leu Leu Glu Ser Val Gln Lys Leu Asp Leu Glu Ala Thr Ile Asp  
 385 390 395 400  
 Lys Val Val Lys Ile Tyr Asp Ala Leu Ile Tyr Met Gln Val Asn Ser  
 405 410 415  
 Ser Phe Glu Asp His Ile Leu Glu Asp Leu Cys Gly Met Leu Ser Leu  
 420 425 430  
 Pro Trp Ile Tyr Ser His Ser Asp Asp Gly Cys Leu Lys Leu Thr Thr  
 435 440 445  
 Phe Ala Ala Asn Leu Leu Thr Leu Ser Cys Arg Ile Ser Asp Ser Tyr  
 450 455 460  
 Ser Pro Gln Ala Gln Ser Arg Cys Val Phe Leu Leu Thr Leu Phe Pro  
 465 470 475 480  
 Arg Arg Ile Phe Leu Glu Trp Arg Thr Ala Val Tyr Asn Trp Ala Leu  
 485 490 495  
 Gln Ser Ser His Glu Val Ile Arg Ala Ser Cys Val Ser Gly Phe Phe  
 500 505 510

-176-

Ile Leu Leu Gln Gln Gln Asn Ser Cys Asn Arg Val Pro Lys Ile Leu  
 515 520 525  
 Ile Asp Lys Val Lys Asp Asp Ser Asp Ile Val Lys Lys Glu Phe Ala  
 530 535 540  
 Ser Ile Leu Gly Gln Leu Val Cys Thr Leu His Gly Met Phe Tyr Leu  
 545 550 555 560  
 Thr Ser Ser Leu Thr Glu Pro Phe Ser Glu His Gly His Val Asp Leu  
 565 570 575  
 Phe Cys Arg Asn Leu Lys Ala Thr Ser Gln His Glu Cys Ser Ser Ser  
 580 585 590  
 Gln Leu Lys Ala Ser Val Cys Lys Pro Phe Leu Phe Leu Leu Lys Lys  
 595 600 605  
 Lys Ile Pro Ser Pro Val Lys Leu Ala Phe Ile Asp Asn Leu His His  
 610 615 620  
 Leu Cys Lys His Leu Asp Phe Arg Glu Asp Glu Thr Asp Val Lys Ala  
 625 630 635 640  
 Val Leu Gly Thr Leu Leu Asn Leu Met Glu Asp Pro Asp Lys Asp Val  
 645 650 655  
 Arg Val Ala Phe Ser Gly Asn Ile Lys His Ile Leu Glu Ser Leu Asp  
 660 665 670  
 Ser Glu Asp Gly Phe Ile Lys Glu Leu Phe Val Leu Arg Met Lys Glu  
 675 680 685  
 Ala Tyr Thr His Ala Gln Ile Ser Arg Asn Asn Glu Leu Lys Asp Thr  
 690 695 700  
 Leu Ile Leu Thr Thr Gly Asp Ile Gly Arg Ala Ala Lys Gly Asp Leu  
 705 710 715 720  
 Val Pro Phe Ala Leu Leu His Leu Leu His Cys Leu Leu Ser Lys Ser  
 725 730 735  
 Ala Ser Val Ser Gly Ala Ala Tyr Thr Glu Ile Arg Ala Leu Val Ala  
 740 745 750  
 Ala Lys Ser Val Lys Leu Gln Ser Phe Phe Ser Gln Tyr Lys Lys Pro  
 755 760 765  
 Ile Cys Gln Phe Leu Val Glu Ser Leu His Ser Ser Gln Met Thr Ala  
 770 775 780  
 Leu Pro Asn Thr Pro Cys Gln Asn Ala Asp Val Arg Lys Gln Asp Val  
 785 790 795 800  
 Ala His Gln Arg Glu Met Ala Leu Asn Thr Leu Ser Glu Ile Ala Asn  
 805 810 815  
 Val Phe Asp Phe Pro Asp Leu Asn Arg Phe Leu Thr Arg Thr Leu Gln  
 820 825 830  
 Val Leu Leu Pro Asp Leu Ala Ala Lys Ala Ser Pro Ala Ala Ser Ala  
 835 840 845  
 Leu Ile Arg Thr Leu Gly Lys Gln Leu Asn Val Asn Arg Arg Glu Ile  
 850 855 860

-177-

Leu Ile Asn Asn Phe Lys Tyr Ile Phe Ser His Leu Val Cys Ser Cys  
 865 870 875 880  
 Ser Lys Asp Glu Leu Glu Arg Ala Leu His Tyr Leu Lys Asn Glu Thr  
 885 890 895  
 Glu Ile Glu Leu Gly Ser Leu Leu Arg Gln Asp Phe Gln Gly Leu His  
 900 905 910  
 Asn Glu Leu Leu Leu Arg Ile Gly Glu His Tyr Gln Gln Val Phe Asn  
 915 920 925  
 Gly Leu Ser Ile Leu Ala Ser Phe Ala Ser Ser Asp Asp Pro Tyr Gln  
 930 935 940  
 Gly Pro Arg Asp Ile Ile Ser Pro Glu Leu Met Ala Asp Tyr Leu Gln  
 945 950 955 960  
 Pro Lys Leu Leu Gly Ile Leu Ala Phe Phe Asn Met Gln Leu Leu Ser  
 965 970 975  
 Ser Ser Val Gly Ile Glu Asp Lys Lys Met Ala Leu Asn Ser Leu Met  
 980 985 990  
 Ser Leu Met Lys Leu Met Gly Pro Lys His Val Ser Ser Val Arg Val  
 995 1000 1005  
 Lys Met Met Thr Thr Leu Arg Thr Gly Leu Arg Phe Lys Asp Asp Phe  
 1010 1015 1020  
 Pro Glu Leu Cys Cys Arg Ala Trp Asp Cys Phe Val Arg Cys Leu Asp  
 1025 1030 1035 1040  
 His Ala Cys Leu Gly Ser Leu Leu Ser His Val Ile Val Ala Leu Leu  
 1045 1050 1055  
 Pro Leu Ile His Ile Gln Pro Lys Glu Thr Ala Ala Ile Phe His Tyr  
 1060 1065 1070  
 Leu Ile Ile Glu Asn Arg Asp Ala Val Gln Asp Phe Leu His Glu Ile  
 1075 1080 1085  
 Tyr Phe Leu Pro Asp His Pro Glu Leu Lys Lys Ile Lys Ala Val Leu  
 1090 1095 1100  
 Gln Glu Tyr Arg Lys Glu Thr Ser Glu Ser Thr Asp Leu Gln Thr Thr  
 1105 1110 1115 1120  
 Leu Gln Leu Ser Met Lys Ala Ile Gln His Glu Asn Val Asp Val Arg  
 1125 1130 1135  
 Ile His Ala Leu Thr Ser Leu Lys Glu Thr Leu Tyr Lys Asn Gln Glu  
 1140 1145 1150  
 Lys Leu Ile Lys Tyr Ala Thr Asp Ser Glu Thr Val Glu Pro Ile Ile  
 1155 1160 1165  
 Ser Gln Leu Val Thr Val Leu Leu Lys Gly Cys Gln Asp Ala Asn Ser  
 1170 1175 1180  
 Gln Ala Arg Leu Leu Cys Gly Glu Cys Leu Gly Glu Leu Gly Ala Ile  
 1185 1190 1195 1200  
 Asp Pro Gly Arg Leu Asp Phe Ser Thr Thr Glu Thr Gln Gly Lys Asp  
 1205 1210 1215

-178-

Phe Thr Phe Val Thr Gly Val Glu Asp Ser Ser Phe Ala Tyr Gly Leu  
 1220 1225 1230  
 Leu Met Glu Leu Thr Arg Ala Tyr Leu Ala Tyr Ala Asp Asn Ser Arg  
 1235 1240 1245  
 Ala Gln Asp Ser Ala Ala Tyr Ala Ile Gln Glu Leu Leu Ser Ile Tyr  
 1250 1255 1260  
 Asp Cys Arg Glu Met Glu Thr Asn Gly Pro Gly His Gln Leu Trp Arg  
 1265 1270 1275 1280  
 Arg Phe Pro Glu His Val Arg Glu Ile Leu Glu Pro His Leu Asn Thr  
 1285 1290 1295  
 Arg Tyr Lys Ser Ser Gln Lys Ser Thr Asp Trp Ser Gly Val Lys Lys  
 1300 1305 1310  
 Pro Ile Tyr Leu Ser Lys Leu Gly Ser Asn Phe Ala Glu Trp Ser Ala  
 1315 1320 1325  
 Ser Trp Ala Gly Tyr Leu Ile Thr Lys Val Arg His Asp Leu Ala Ser  
 1330 1335 1340  
 Lys Ile Phe Thr Cys Cys Ser Ile Met Met Lys His Asp Phe Lys Val  
 1345 1350 1355 1360  
 Thr Ile Tyr Leu Leu Pro His Ile Leu Val Tyr Val Leu Leu Gly Cys  
 1365 1370 1375  
 Asn Gln Glu Asp Gln Gln Glu Val Tyr Ala Glu Ile Met Ala Val Leu  
 1380 1385 1390  
 Lys His Asp Asp Gln His Thr Ile Asn Thr Gln Asp Ile Ala Ser Asp  
 1395 1400 1405  
 Leu Cys Gln Leu Ser Thr Gln Thr Val Phe Ser Met Leu Asp His Leu  
 1410 1415 1420  
 Thr Gln Trp Ala Arg His Lys Phe Gln Ala Leu Lys Ala Glu Lys Cys  
 1425 1430 1435 1440  
 Pro His Ser Lys Ser Asn Arg Asn Lys Val Asp Ser Met Val Ser Thr  
 1445 1450 1455  
 Val Asp Tyr Glu Asp Tyr Gln Ser Val Thr Arg Phe Leu Asp Leu Ile  
 1460 1465 1470  
 Pro Gln Asp Thr Leu Ala Val Ala Ser Phe Arg Ser Lys Ala Tyr Thr  
 1475 1480 1485  
 Arg Ala Val Met His Phe Glu Ser Phe Ile Thr Glu Lys Lys Gln Asn  
 1490 1495 1500  
 Ile Gln Glu His Leu Gly Phe Leu Gln Lys Leu Tyr Ala Ala Met His  
 1505 1510 1515 1520  
 Glu Pro Asp Gly Val Ala Gly Val Ser Ala Ile Arg Lys Ala Glu Pro  
 1525 1530 1535  
 Ser Leu Lys Glu Gln Ile Leu Glu His Glu Ser Leu Gly Leu Leu Arg  
 1540 1545 1550  
 Asp Ala Thr Ala Cys Tyr Asp Arg Ala Ile Gln Leu Glu Pro Asp Gln  
 1555 1560 1565



-179-

Ile Ile His Tyr His Gly Val Val Lys Ser Met Leu Gly Leu Gly Gln  
 1570 1575 1580  
 Leu Ser Thr Val Ile Thr Gln Val Asn Gly Val His Ala Asn Arg Ser  
 1585 1590 1595 1600  
 Glu Trp Thr Asp Glu Leu Asn Thr Tyr Arg Val Glu Ala Ala Trp Lys  
 1605 1610 1615  
 Leu Ser Gln Trp Asp Leu Val Glu Asn Tyr Leu Ala Ala Asp Gly Lys  
 1620 1625 1630  
 Ser Thr Thr Trp Ser Val Arg Leu Gly Gln Leu Leu Leu Ser Ala Lys  
 1635 1640 1645  
 Lys Arg Asp Ile Thr Ala Phe Tyr Asp Ser Leu Lys Leu Val Arg Ala  
 1650 1655 1660  
 Glu Gln Ile Val Pro Leu Ser Ala Ala Ser Phe Glu Arg Gly Ser Tyr  
 1665 1670 1675 1680  
 Gln Arg Gly Tyr Glu Tyr Ile Val Arg Leu His Met Leu Cys Glu Leu  
 1685 1690 1695  
 Glu His Ser Ile Lys Pro Leu Phe Gln His Ser Pro Gly Asp Ser Ser  
 1700 1705 1710  
 Gln Glu Asp Ser Leu Asn Trp Val Ala Arg Leu Glu Met Thr Gln Asn  
 1715 1720 1725  
 Ser Tyr Arg Ala Lys Glu Pro Ile Leu Ala Leu Arg Arg Ala Leu Leu  
 1730 1735 1740  
 Ser Leu Asn Lys Arg Pro Asp Tyr Asn Glu Met Val Gly Glu Cys Trp  
 1745 1750 1755 1760  
 Leu Gln Ser Ala Arg Val Ala Arg Lys Ala Gly His His Gln Thr Ala  
 1765 1770 1775  
 Tyr Asn Ala Leu Leu Asn Ala Gly Glu Ser Arg Leu Ala Glu Leu Tyr  
 1780 1785 1790  
 Val Glu Arg Ala Lys Trp Leu Trp Ser Lys Gly Asp Val His Gln Ala  
 1795 1800 1805  
 Leu Ile Val Leu Gln Lys Gly Val Glu Leu Cys Phe Pro Glu Asn Glu  
 1810 1815 1820  
 Thr Pro Pro Glu Gly Lys Asn Met Leu Ile His Gly Arg Ala Met Leu  
 1825 1830 1835 1840  
 Leu Val Gly Arg Phe Met Glu Glu Thr Ala Asn Phe Glu Ser Asn Ala  
 1845 1850 1855  
 Ile Met Lys Lys Tyr Lys Asp Val Thr Ala Cys Leu Pro Glu Trp Glu  
 1860 1865 1870  
 Asp Gly His Phe Tyr Leu Ala Lys Tyr Tyr Asp Lys Leu Met Pro Met  
 1875 1880 1885  
 Val Thr Asp Asn Lys Met Glu Lys Gln Gly Asp Leu Ile Arg Tyr Ile  
 1890 1895 1900  
 Val Leu His Phe Gly Arg Ser Leu Gln Tyr Gly Asn Gln Phe Ile Tyr  
 1905 1910 1915 1920

-180-

Gln Ser Met Pro Arg Met Leu Thr Leu Trp Leu Asp Tyr Gly Thr Lys  
 1925 1930 1935  
 Ala Tyr Glu Trp Glu Lys Ala Gly Arg Ser Asp Arg Val Gln Met Arg  
 1940 1945 1950  
 Asn Asp Leu Gly Lys Ile Asn Lys Val Ile Thr Glu His Thr Asn Tyr  
 1955 1960 1965  
 Leu Ala Pro Tyr Gln Phe Leu Thr Ala Phe Ser Gln Leu Ile Ser Arg  
 1970 1975 1980  
 Ile Cys His Ser His Asp Glu Val Phe Val Val Leu Met Glu Ile Ile  
 1985 1990 1995 2000  
 Ala Lys Val Phe Leu Ala Tyr Pro Gln Gln Ala Met Trp Met Met Thr  
 2005 2010 2015  
 Ala Val Ser Lys Ser Ser Tyr Pro Met Arg Val Asn Arg Cys Lys Glu  
 2020 2025 2030  
 Ile Leu Asn Lys Ala Ile His Met Lys Lys Ser Leu Glu Lys Phe Val  
 2035 2040 2045  
 Gly Asp Ala Thr Arg Leu Thr Asp Lys Leu Leu Glu Leu Cys Asn Lys  
 2050 2055 2060  
 Pro Val Asp Gly Ser Ser Ser Thr Leu Ser Met Ser Thr His Phe Lys  
 2065 2070 2075 2080  
 Met Leu Lys Lys Leu Val Glu Glu Ala Thr Phe Ser Glu Ile Leu Ile  
 2085 2090 2095  
 Pro Leu Gln Ser Val Met Ile Pro Thr Leu Pro Ser Ile Leu Gly Thr  
 2100 2105 2110  
 His Ala Asn His Ala Ser His Glu Pro Phe Pro Gly His Trp Ala Tyr  
 2115 2120 2125  
 Ile Ala Gly Phe Asp Asp Met Val Glu Ile Leu Ala Ser Leu Gln Lys  
 2130 2135 2140  
 Pro Lys Lys Ile Ser Leu Lys Gly Ser Asp Gly Lys Phe Tyr Ile Met  
 2145 2150 2155 2160  
 Met Cys Lys Pro Lys Asp Asp Leu Arg Lys Asp Cys Arg Leu Met Glu  
 2165 2170 2175  
 Phe Asn Ser Leu Ile Asn Lys Cys Leu Arg Lys Asp Ala Glu Ser Arg  
 2180 2185 2190  
 Arg Arg Glu Leu His Ile Arg Thr Tyr Ala Val Ile Pro Leu Asn Asp  
 2195 2200 2205  
 Glu Cys Gly Ile Ile Glu Trp Val Asn Asn Thr Ala Gly Leu Arg Pro  
 2210 2215 2220  
 Ile Leu Thr Lys Leu Tyr Lys Glu Lys Gly Val Tyr Met Thr Gly Lys  
 2225 2230 2235 2240  
 Glu Leu Arg Gln Cys Met Leu Pro Lys Ser Ala Ala Leu Ser Glu Lys  
 2245 2250 2255  
 Leu Lys Val Phe Arg Glu Phe Leu Leu Pro Arg His Pro Pro Ile Phe  
 2260 2265 2270

-181-

His Glu Trp Phe Leu Arg Thr Phe Pro Asp Pro Thr Ser Trp Tyr Ser  
 2275 2280 2285  
 Ser Arg Ser Ala Tyr Cys Arg Ser Thr Ala Val Met Ser Met Val Gly  
 2290 2295 2300  
 Tyr Ile Leu Gly Leu Gly Asp Arg His Gly Glu Asn Ile Leu Phe Asp  
 2305 2310 2315 2320  
 Ser Leu Thr Gly Glu Cys Val His Val Asp Phe Asn Cys Leu Phe Asn  
 2325 2330 2335  
 Lys Gly Glu Thr Phe Glu Val Pro Glu Ile Val Pro Phe Arg Leu Thr  
 2340 2345 2350  
 His Asn Met Val Asn Gly Met Gly Pro Met Gly Thr Glu Gly Leu Phe  
 2355 2360 2365  
 Arg Arg Ala Cys Glu Val Thr Met Arg Leu Met Arg Asp Gln Arg Glu  
 2370 2375 2380  
 Pro Leu Met Ser Val Leu Lys Thr Phe Leu His Asp Pro Leu Val Glu  
 2385 2390 2395 2400  
 Trp Ser Lys Pro Val Lys Gly His Ser Lys Ala Pro Leu Asn Glu Thr  
 2405 2410 2415  
 Gly Glu Val Val Asn Glu Lys Ala Lys Thr His Val Leu Asp Ile Glu  
 2420 2425 2430  
 Gln Arg Leu Gln Gly Val Ile Lys Thr Arg Asn Arg Val Thr Gly Leu  
 2435 2440 2445  
 Pro Leu Ser Ile Glu Gly His Val His Tyr Leu Ile Gln Glu Ala Thr  
 2450 2455 2460  
 Asp Glu Asn Leu Leu Cys Gln Met Tyr Leu Gly Trp Thr Pro Tyr Met  
 2465 2470 2475 2480

## (2) INFORMATION FOR SEQ ID NO:34:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 9385 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (v) FRAGMENT TYPE: linear

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 190..9357

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

GCGAGAGGAG TCGGGATCTG CGCTGCAGCC ACCGCCGCGG TTGATACTAC TTTGACCTTC	60
CGAGTGCAGT GAGGCATACA TCACAATTG GAATTATGCA TTGGTTTATC AATTACTTG	120
TTTATTGTCA CCCTGCTGCC CAGATATGAC TTCATGAGGA CAGTGATGTG TGTCTGAAA	180

-182-

TTGTGAACC	ATG	AGT	CTA	GTA	CTT	AAT	GAT	CTG	CTT	ATC	TGC	TGC	CGT		228	
	Met	Ser	Leu	Val	Leu	Asn	Asp	Leu	Leu	Ile	Cys	Cys	Arg			
	1				5					10						
CAA	CTA	GAA	CAT	GAT	AGA	GCT	ACA	GAA	CGA	AAG	AAA	GAA	GTT	GAG	AAA	276
Gln	Leu	Glu	His	Asp	Arg	Ala	Thr	Glu	Arg	Lys	Lys	Glu	Val	Glu	Lys	
	15					20					25					
TTT	AAG	CGC	CTG	ATT	CGA	GAT	CCT	GAA	ACA	ATT	AAA	CAT	CTA	GAT	CGG	324
Phe	Lys	Arg	Leu	Ile	Arg	Asp	Pro	Glu	Thr	Ile	Lys	His	Leu	Asp	Arg	
	30				35					40					45	
CAT	TCA	GAT	TCC	AAA	CAA	GGA	AAA	TAT	TTG	AAT	TGG	GAT	GCT	GTT	TTT	372
His	Ser	Asp	Ser	Lys	Gln	Gly	Lys	Tyr	Leu	Asn	Trp	Asp	Ala	Val	Phe	
				50					55					60		
AGA	TTT	TTA	CAG	AAA	TAT	ATT	CAG	AAA	GAA	ACA	GAA	TGT	CTG	AGA	ATA	420
Arg	Phe	Leu	Gln	Lys	Tyr	Ile	Gln	Lys	Glu	Thr	Glu	Cys	Leu	Arg	Ile	
			65					70					75			
GCA	AAA	CCA	AAT	GTA	TCA	GCC	TCA	ACA	CAA	GCC	TCC	AGG	CAG	AAA	AAG	468
Ala	Lys	Pro	Asn	Val	Ser	Ala	Ser	Thr	Gln	Ala	Ser	Arg	Gln	Lys	Lys	
		80					85					90				
ATG	CAG	GAA	ATC	AGT	AGT	TTG	GTC	AAA	TAC	TTC	ATC	AAA	TGT	GCA	AAC	516
Met	Gln	Glu	Ile	Ser	Ser	Leu	Val	Lys	Tyr	Phe	Ile	Lys	Cys	Ala	Asn	
	95					100					105					
AGA	AGA	GCA	CCT	AGG	CTA	AAA	TGT	CAA	GAA	CTC	TTA	AAT	TAT	ATC	ATG	564
Arg	Arg	Ala	Pro	Arg	Leu	Lys	Cys	Gln	Glu	Leu	Leu	Asn	Tyr	Ile	Met	
	110				115					120					125	
GAT	ACA	GTG	AAA	GAT	TCA	TCT	AAT	GGT	GCT	ATT	TAC	GGA	GCT	GAT	TGT	612
Asp	Thr	Val	Lys	Asp	Ser	Ser	Asn	Gly	Ala	Ile	Tyr	Gly	Ala	Asp	Cys	
				130					135					140		
AGC	AAC	ATA	CTA	CTC	AAA	GAC	ATT	CTT	TCT	GTG	AGA	AAA	TAC	TGG	TGT	660
Ser	Asn	Ile	Leu	Lys	Asp	Ile	Leu	Ser	Val	Arg	Lys	Tyr	Trp	Cys		
			145				150						155			
GAA	ATA	TCT	CAG	CAA	CAG	TGG	TTA	GAA	TTG	TTC	TCT	GTG	TAC	TTC	AGG	708
Glu	Ile	Ser	Gln	Gln	Gln	Trp	Leu	Glu	Leu	Phe	Ser	Val	Tyr	Phe	Arg	
	160					165						170				
CTC	TAT	CTG	AAA	CCT	TCA	CAA	GAT	GTT	CAT	AGA	GTT	TTA	GTG	GCT	AGA	756
Leu	Tyr	Leu	Lys	Pro	Ser	Gln	Asp	Val	His	Arg	Val	Leu	Val	Ala	Arg	
	175					180					185					
ATA	ATT	CAT	GCT	GTT	ACC	AAA	GGA	TGC	TGT	TCT	CAG	ACT	GAC	GGA	TTA	804
Ile	Ile	His	Ala	Val	Thr	Lys	Gly	Cys	Cys	Ser	Gln	Thr	Asp	Gly	Leu	
	190				195					200					205	
AAT	TCC	AAA	TTT	TTG	GAC	TTT	TTT	TCC	AAG	GCT	ATT	CAG	TGT	GCG	AGA	852
Asn	Ser	Lys	Phe	Leu	Asp	Phe	Phe	Ser	Lys	Ala	Ile	Gln	Cys	Ala	Arg	
				210					215					220		
CAA	GAA	AAG	AGC	TCT	TCA	GGT	CTA	AAT	CAT	ATC	TTA	GCA	GCT	CTT	ACT	900
Gln	Glu	Lys	Ser	Ser	Ser	Gly	Leu	Asn	His	Ile	Leu	Ala	Ala	Leu	Thr	
			225					230					235			
ATC	TTC	CTC	AAG	ACT	TTG	GCT	GTC	AAC	TTT	CGA	ATT	CGA	GTG	TGT	GAA	948
Ile	Phe	Leu	Lys	Thr	Leu	Ala	Val	Asn	Phe	Arg	Ile	Arg	Val	Cys	Glu	
		240				245						250				
TTA	GGA	GAT	GAA	ATT	CTT	CCC	ACT	TTG	CTT	TAT	ATT	TGG	ACT	CAA	CAT	996
Leu	Gly	Asp	Glu	Ile	Leu	Pro	Thr	Leu	Leu	Tyr	Ile	Trp	Thr	Gln	His	
	255					260					265					

-183-

AGG CTT AAT GAT TCT TTA AAA GAA GTC ATT ATT GAA TTA TTT CAA CTG	1044
Arg Leu Asn Asp Ser Leu Lys Glu Val Ile Ile Glu Leu Phe Gln Leu	
270 275 280 285	
CAA ATT TAT ATC CAT CAT CCG AAA GGA GCC AAA ACC CAA GAA AAA GGT	1092
Gln Ile Tyr Ile His His Pro Lys Gly Ala Lys Thr Gln Glu Lys Gly	
290 295 300	
GCT TAT GAA TCA ACA AAA TGG AGA AGT ATT TTA TAC AAC TTA TAT GAT	1140
Ala Tyr Glu Ser Thr Lys Trp Arg Ser Ile Leu Tyr Asn Leu Tyr Asp	
305 310 315	
CTG CTA GTG AAT GAG ATA AGT CAT ATA GGA AGT AGA GGA AAG TAT TCT	1188
Leu Leu Val Asn Glu Ile Ser His Ile Gly Ser Arg Gly Lys Tyr Ser	
320 325 330	
TCA GGA TTT CGT AAT ATT GCC GTC AAA GAA AAT TTG ATT GAA TTG ATG	1236
Ser Gly Phe Arg Asn Ile Ala Val Lys Glu Asn Leu Ile Glu Leu Met	
335 340 345	
GCA GAT ATC TGT CAC CAG GTT TTT AAT GAA GAT ACC AGA TCC TTG GAG	1284
Ala Asp Ile Cys His Gln Val Phe Asn Glu Asp Thr Arg Ser Leu Glu	
350 355 360 365	
ATT TCT CAA TCT TAC ACT ACT ACA CAA AGA GAA TCT AGT GAT TAC AGT	1332
Ile Ser Gln Ser Tyr Thr Thr Thr Gln Arg Glu Ser Ser Asp Tyr Ser	
370 375 380	
GTC CCT TGC AAA AGG AAG AAA ATA GAA CTA GGC TGG GAA GTA ATA AAA	1380
Val Pro Cys Lys Arg Lys Lys Ile Glu Leu Gly Trp Glu Val Ile Lys	
385 390 395	
GAT CAC CTT CAG AAG TCA CAG AAT GAT TTT GAT CTT GTG CCT TGG CTA	1428
Asp His Leu Gln Lys Ser Gln Asn Asp Phe Asp Leu Val Pro Trp Leu	
400 405 410	
CAG ATT GCA ACC CAA TTA ATA TCA AAG TAT CCT GCA AGT TTA CCT AAC	1476
Gln Ile Ala Thr Gln Leu Ile Ser Lys Tyr Pro Ala Ser Leu Pro Asn	
415 420 425	
TGT GAG CTG TCT CCA TTA CTG ATG ATA CTA TCT CAG CTT CTA CCC CAA	1524
Cys Glu Leu Ser Pro Leu Leu Met Ile Leu Ser Gln Leu Leu Pro Gln	
430 435 440 445	
CAG CGA CAT GGG GAA CGT ACA CCA TAT GTG TTA CGA TGC CTT ACG GAA	1572
Gln Arg His Gly Glu Arg Thr Pro Tyr Val Leu Arg Cys Leu Thr Glu	
450 455 460	
GTT GCA TTG TGT CAA GAC AAG AGG TCA AAC CTA GAA AGC TCA CAA AAG	1620
Val Ala Leu Cys Gln Asp Lys Arg Ser Asn Leu Glu Ser Gln Lys	
465 470 475	
TCA GAT TTA TTA AAA CTC TGG AAT AAA ATT TGG TGT ATT ACC TTT CGT	1668
Ser Asp Leu Leu Lys Leu Trp Asn Lys Ile Trp Cys Ile Thr Phe Arg	
480 485 490	
GGT ATA AGT TCT GAG CAA ATA CAA GCT GAA AAC TTT GGC TTA CTT GGA	1716
Gly Ile Ser Ser Glu Gln Ile Gln Ala Glu Asn Phe Gly Leu Leu Gly	
495 500 505	
GCC ATA ATT CAG GGT AGT TTA GTT GAG GTT GAC AGA GAA TTC TGG AAG	1764
Ala Ile Ile Gln Gly Ser Leu Val Glu Val Asp Arg Glu Phe Trp Lys	
510 515 520 525	
TTA TTT ACT GGG TCA GCC TGC AGA CCT TCA TGT CCT GCA GTA TGC TGT	1812
Leu Phe Thr Gly Ser Ala Cys Arg Pro Ser Cys Pro Ala Val Cys Cys	
530 535 540	

-184-

TTG	ACT	TTG	GCA	CTG	ACC	ACC	AGT	ATA	GTT	CCA	GGA	GCG	GTA	AAA	ATG	1860
Leu	Thr	Leu	Ala	Leu	Thr	Thr	Ser	Ile	Val	Pro	Gly	Ala	Val	Lys	Met	
			545					550					555			
GGA	ATA	GAG	CAA	AAT	ATG	TGT	GAA	GTA	AAT	AGA	AGC	TTT	TCT	TTA	AAG	1908
Gly	Ile	Glu	Gln	Asn	Met	Cys	Glu	Val	Asn	Arg	Ser	Phe	Ser	Leu	Lys	
		560					565					570				
GAA	TCA	ATA	ATG	AAA	TGG	CTC	TTA	TTC	TAT	CAG	TTA	GAG	GGT	GAC	TTA	1956
Glu	Ser	Ile	Met	Lys	Trp	Leu	Leu	Phe	Tyr	Gln	Leu	Glu	Gly	Asp	Leu	
	575					580					585					
GAA	AAT	AGC	ACA	GAA	GTG	CCT	CCA	ATT	CTT	CAC	AGT	AAT	TTT	CCT	CAT	2004
Glu	Asn	Ser	Thr	Glu	Val	Pro	Pro	Ile	Leu	His	Ser	Asn	Phe	Pro	His	
590					595					600					605	
CTT	GTA	CTG	GAG	AAA	ATT	CTT	GTG	AGT	CTC	ACT	ATG	AAA	AAC	TGT	AAA	2052
Leu	Val	Leu	Glu	Lys	Ile	Leu	Val	Ser	Leu	Thr	Met	Lys	Asn	Cys	Lys	
				610					615					620		
GCT	GCA	ATG	AAT	TTT	TTC	CAA	AGC	GTG	CCA	GAA	TGT	GAA	CAC	CAC	CAA	2100
Ala	Ala	Met	Asn	Phe	Phe	Gln	Ser	Val	Pro	Glu	Cys	Glu	His	His	Gln	
			625					630					635			
AAA	GAT	AAA	GAA	GAA	CTT	TCA	TTC	TCA	GAA	GTA	GAA	GAA	CTA	TTT	CTT	2148
Lys	Asp	Lys	Glu	Glu	Leu	Ser	Phe	Ser	Glu	Val	Glu	Glu	Leu	Phe	Leu	
		640					645					650				
CAG	ACA	ACT	TTT	GAC	AAG	ATG	GAC	TTT	TTA	ACC	ATT	GTG	AGA	GAA	TGT	2196
Gln	Thr	Thr	Phe	Asp	Lys	Met	Asp	Phe	Leu	Thr	Ile	Val	Arg	Glu	Cys	
	655					660					665					
GGT	ATA	GAA	AAG	CAC	CAG	TCC	AGT	ATT	GGC	TTC	TCT	GTC	CAC	CAG	AAT	2244
Gly	Ile	Glu	Lys	His	Gln	Ser	Ser	Ile	Gly	Phe	Ser	Val	His	Gln	Asn	
670					675				680						685	
CTC	AAG	GAA	TCA	CTG	GAT	CGC	TGT	CTT	CTG	GGA	TTA	TCA	GAA	CAG	CTT	2292
Leu	Lys	Glu	Ser	Leu	Asp	Arg	Cys	Leu	Leu	Gly	Leu	Ser	Glu	Gln	Leu	
				690					695					700		
CTG	AAT	AAT	TAC	TCA	TCT	GAG	ATT	ACA	AAT	TCA	GAA	ACT	CTT	GTC	CGG	2340
Leu	Asn	Asn	Tyr	Ser	Ser	Glu	Ile	Thr	Asn	Ser	Glu	Thr	Leu	Val	Arg	
			705					710					715			
TGT	TCA	CGT	CTT	TTG	GTG	GGT	GTC	CTT	GGC	TGC	TAC	TGT	TAC	ATG	GGT	2388
Cys	Ser	Arg	Leu	Leu	Val	Gly	Val	Leu	Gly	Cys	Tyr	Cys	Tyr	Met	Gly	
		720					725					730				
GTA	ATA	GCT	GAA	GAG	GAA	GCA	TAT	AAG	TCA	GAA	TTA	TTC	CAG	AAA	GCC	2436
Val	Ile	Ala	Glu	Glu	Glu	Ala	Tyr	Lys	Ser	Glu	Leu	Phe	Gln	Lys	Ala	
	735					740					745					
AAC	TCT	CTA	ATG	CAA	TGT	GCA	GGA	GAA	AGT	ATC	ACT	CTG	TTT	AAA	AAT	2484
Asn	Ser	Leu	Met	Gln	Cys	Ala	Gly	Glu	Ser	Ile	Thr	Leu	Phe	Lys	Asn	
750					755				760						765	
AAG	ACA	AAT	GAG	GAA	TTC	AGA	ATT	GGT	TCC	TTG	AGA	AAT	ATG	ATG	CAG	2532
Lys	Thr	Asn	Glu	Glu	Phe	Arg	Ile	Gly	Ser	Leu	Arg	Asn	Met	Met	Gln	
				770					775					780		
CTA	TGT	ACA	CGT	TGC	TTG	AGC	AAC	TGT	ACC	AAG	AAG	AGT	CCA	AAT	AAG	2580
Leu	Cys	Thr	Arg	Cys	Leu	Ser	Asn	Cys	Thr	Lys	Lys	Ser	Pro	Asn	Lys	
			785					790					795			
ATT	GCA	TCT	GGC	TTT	TTC	CTG	CGA	TTG	TTA	ACA	TCA	AAG	CTA	ATG	AAT	2628
Ile	Ala	Ser	Gly	Phe	Phe	Leu	Arg	Leu	Leu	Thr	Ser	Lys	Leu	Met	Asn	
		800					805					810				

-185-

GAC Asp 815	ATT Ile	GCA Ala	GAT Asp	ATT Ile	TGT Cys	AAA Lys	AGT Ser	TTA Leu	GCA Ala	TCC Ser	TTC Phe	ATC Ile	AAA Lys	AAG Lys	CCA Pro	2676
TTT Phe 830	GAC Asp	CGT Arg	GGA Gly	GAA Glu	GTA Val	GAA Glu	TCA Ser	ATG Met	GAA Glu	GAT Asp	GAT Asp	ACT Thr	AAT Asn	GGA Gly	AAT Asn 845	2724
CTA Leu	ATG Met	GAG Glu	GTG Val	GAG Glu	GAT Asp	CAG Gln	TCA Ser	TCC Ser	ATG Met	AAT Asn	CTA Leu	TTT Phe	AAC Asn	GAT Asp	TAC Tyr 860	2772
CCT Pro	GAT Asp	AGT Ser	AGT Ser	GTT Val	AGT Ser	GAT Asp	GCA Ala	AAC Asn	GAA Glu	CCT Pro	GGA Gly	GAG Glu	AGC Ser	CAA Gln	AGT Ser	2820
ACC Thr	ATA Ile	GGT Gly	GCC Ala	ATT Ile	AAT Asn	CCT Pro	TTA Leu	GCT Ala	GAA Glu	GAA Glu	TAT Tyr	CTG Leu	TCA Ser	AAG Lys	CAA Gln	2868
GAT Asp 895	CTA Leu	CTT Leu	TTC Phe	TTA Leu	GAC Asp	ATG Met	CTC Leu	AAG Lys	TTC Phe	TTG Leu	TGT Cys	TTG Leu	TGT Cys	GTA Val	ACT Thr	2916
ACT Thr 910	GCT Ala	CAG Gln	ACC Thr	AAT Asn	ACT Thr	GTG Val	TCC Ser	TTT Phe	AGG Arg	GCA Ala	GCT Ala	GAT Asp	ATT Ile	CGG Arg	AGG Arg 925	2964
AAA Lys	TTG Leu	TTA Leu	ATG Met	TTA Leu	ATT Ile	GAT Asp	TCT Ser	AGC Ser	ACG Thr	CTA Leu	GAA Glu	CCT Pro	ACC Thr	AAA Lys	TCC Ser 940	3012
CTC Leu	CAC His	CTG Leu	CAT His	ATG Met	TAT Tyr	CTA Leu	ATG Met	CTT Leu	TTA Leu	AAG Lys	GAG Glu	CTT Leu	CCT Pro	GGA Gly	GAA Glu	3060
GAG Glu	TAC Tyr	CCC Pro	TTG Leu	CCA Pro	ATG Met	GAA Glu	GAT Asp	GTT Val	CTT Leu	GAA Glu	CTT Leu	CTG Leu	AAA Lys	CCA Pro	CTA Leu	3108
TCC Ser 975	AAT Asn	GTG Val	TGT Cys	TCT Ser	TTG Leu	TAT Tyr	CGT Arg	CGT Arg	GAC Asp	CAA Gln	GAT Asp	GTT Val	TGT Cys	AAA Lys	ACT Thr	3156
ATT Ile 990	TTA Leu	AAC Asn	CAT His	GTC Val	CTT Leu	CAT His	GTA Val	GTG Val	AAA Lys	AAC Asn	CTA Leu	GGT Gly	CAA Gln	AGC Ser	AAT Asn 1005	3204
ATG Met	GAC Asp	TCT Ser	GAG Glu	AAC Asn	ACA Thr	AGG Arg	GAT Asp	GCT Ala	CAA Gln	GGA Gly	CAG Gln	TTT Phe	CTT Leu	ACA Thr	GTA Val 1020	3252
ATT Ile	GGA Gly	GCA Ala	TTT Phe	TGG Trp	CAT His	CTA Leu	ACA Thr	AAG Lys	GAG Glu	AGG Arg	AAA Lys	TAT Tyr	ATA Ile	TTC Phe	TCT Ser	3300
GTA Val	AGA Arg	ATG Met	GCC Ala	CTA Leu	GTA Val	AAT Asn	TGC Cys	CTT Leu	AAA Lys	ACT Thr	TTG Leu	CTT Leu	GAG Glu	GCT Ala	GAT Asp	3348
CCT Pro	TAT Tyr	TCA Ser	AAA Lys	TGG Trp	GCC Ala	ATT Ile	CTT Leu	AAT Asn	GTA Val	ATG Met	GGA Gly	AAA Lys	GAC Asp	TTT Phe	CCT Pro	3396
GTA Val 1070	AAT Asn	GAA Glu	GTA Val	TTT Phe	ACA Thr	CAA Gln	TTT Phe	CTT Leu	GCT Ala	GAC Asp	AAT Asn	CAT His	CAC His	CAA Gln	GTT Val 1085	3444

-186-

CGC ATG TTG GCT GCA GAG TCA ATC AAT AGA TTG TTC CAG GAC ACG AAG	3492
Arg Met Leu Ala Ala Glu Ser Ile Asn Arg Leu Phe Gln Asp Thr Lys	
1090 1095 1100	
GGA GAT TCT TCC AGG TTA CTG AAA GCA CTT CCT TTG AAG CTT CAG CAA	3540
Gly Asp Ser Arg Leu Leu Lys Ala Leu Pro Leu Lys Leu Gln Gln	
1105 1110 1115	
ACA GCT TTT GAA AAT GCA TAC TTG AAA GCT CAG GAA GGA ATG AGA GAA	3588
Thr Ala Phe Glu Asn Ala Tyr Leu Lys Ala Gln Glu Gly Met Arg Glu	
1120 1125 1130	
ATG TCC CAT AGT GCT GAG AAC CCT GAA ACT TTG GAT GAA ATT TAT AAT	3636
Met Ser His Ser Ala Glu Asn Pro Glu Thr Leu Asp Glu Ile Tyr Asn	
1135 1140 1145	
AGA AAA TCT GTT TTA CTG ACG TTG ATA GCT GTG GTT TTA TCC TGT AGC	3684
Arg Lys Ser Val Leu Leu Thr Leu Ile Ala Val Val Leu Ser Cys Ser	
1150 1155 1160 1165	
CCT ATC TGC GAA AAA CAG GCT TTG TTT GCC CTG TGT AAA TCT GTG AAA	3732
Pro Ile Cys Glu Lys Gln Ala Leu Phe Ala Leu Cys Lys Ser Val Lys	
1170 1175 1180	
GAG AAT GGA TTA GAA CCT CAC CTT GTG AAA AAG GTT TTA GAG AAA GTT	3780
Glu Asn Gly Leu Glu Pro His Leu Val Lys Lys Val Leu Glu Lys Val	
1185 1190 1195	
TCT GAA ACT TTT GGA TAT AGA CGT TTA GAA GAC TTT ATG GCA TCT CAT	3828
Ser Glu Thr Phe Gly Tyr Arg Arg Leu Glu Asp Phe Met Ala Ser His	
1200 1205 1210	
TTA GAT TAT CTG GTT TTG GAA TGG CTA AAT CTT CAA GAT ACT GAA TAC	3876
Leu Asp Tyr Leu Val Leu Glu Trp Leu Asn Leu Gln Asp Thr Glu Tyr	
1215 1220 1225	
AAC TTA TCT TCT TTT CCT TTT ATT TTA TTA AAC TAC ACA AAT ATT GAG	3924
Asn Leu Ser Ser Phe Pro Phe Ile Leu Leu Asn Tyr Thr Asn Ile Glu	
1230 1235 1240 1245	
GAT TTC TAT AGA TCT TGT TAT AAG GTT TTG ATT CCA CAT CTG GTG ATT	3972
Asp Phe Tyr Arg Ser Cys Tyr Lys Val Leu Ile Pro His Leu Val Ile	
1250 1255 1260	
AGA AGT CAT TTT GAT GAG GTG AAG TCC ATT GCT AAT CAG ATT CAA GAG	4020
Arg Ser His Phe Asp Glu Val Lys Ser Ile Ala Asn Gln Ile Gln Glu	
1265 1270 1275	
GAC TGG AAA AGT CTT CTA ACA GAC TGC TTT CCA AAG ATT CTT GTA AAT	4068
Asp Trp Lys Ser Leu Leu Thr Asp Cys Phe Pro Lys Ile Leu Val Asn	
1280 1285 1290	
ATT CTT CCT TAT TTT GCC TAT GAG GGT ACC AGA GAC AGT GGG ATG GCA	4116
Ile Leu Pro Tyr Phe Ala Tyr Glu Gly Thr Arg Asp Ser Gly Met Ala	
1295 1300 1305	
CAG CAA AGA GAG ACT GCT ACC AAG GTC TAT GAT ATG CTT AAA AGT GAA	4164
Gln Gln Arg Glu Thr Ala Thr Lys Val Tyr Asp Met Leu Lys Ser Glu	
1310 1315 1320 1325	
AAC TTA TTG GGA AAA CAG ATT GAT CAC TTA TTC ATT AGT AAT TTA CCA	4212
Asn Leu Leu Gly Lys Gln Ile Asp His Leu Phe Ile Ser Asn Leu Pro	
1330 1335 1340	
GAG ATT GTG GTG GAG TTA TTG ATG ACG TTA CAT GAG CCA GCA AAT TCT	4260
Glu Ile Val Val Glu Leu Leu Met Thr Leu His Glu Pro Ala Asn Ser	
1345 1350 1355	



-187-

AGT GCC AGT CAG AGC ACT GAC CTC TGT GAC TTT TCA GGG GAT TTG GAT Ser Ala Ser Gln Ser Thr Asp Leu Cys Asp Phe Ser Gly Asp Leu Asp 1360 1365 1370	4308
CCT GCT CCT AAT CCA CCT CAT TTT CCA TCG CAT GTG ATT AAA GCA ACA Pro Ala Pro Asn Pro Pro His Phe Pro Ser His Val Ile Lys Ala Thr 1375 1380 1385	4356
TTT GCC TAT ATC AGC AAT TGT CAT AAA ACC AAG TTA AAA AGC ATT TTA Phe Ala Tyr Ile Ser Asn Cys His Lys Thr Lys Leu Lys Ser Ile Leu 1390 1395 1400 1405	4404
GAA ATT CTT TCC AAA AGC CCT GAT TCC TAT CAG AAA ATT CTT CTT GCC Glu Ile Leu Ser Lys Ser Pro Asp Ser Tyr Gln Lys Ile Leu Leu Ala 1410 1415 1420	4452
ATA TGT GAG CAA GCA GCT GAA ACA AAT AAT GTT TAT AAG AAG CAC AGA Ile Cys Glu Gln Ala Ala Glu Thr Asn Asn Val Tyr Lys Lys His Arg 1425 1430 1435	4500
ATT CTT AAA ATA TAT CAC CTG TTT GTT AGT TTA TTA CTG AAA GAT ATA Ile Leu Lys Ile Tyr His Leu Phe Val Ser Leu Leu Leu Lys Asp Ile 1440 1445 1450	4548
AAA AGT GGC TTA GGA GGA GCT TGG GCC TTT GTT CTT CGA GAC GTT ATT Lys Ser Gly Leu Gly Gly Ala Trp Ala Phe Val Leu Arg Asp Val Ile 1455 1460 1465	4596
TAT ACT TTG ATT CAC TAT ATC AAC CAA AGG CCT TCT TGT ATC ATG GAT Tyr Thr Leu Ile His Tyr Ile Asn Gln Arg Pro Ser Cys Ile Met Asp 1470 1475 1480 1485	4644
GTG TCA TTA CGT AGC TTC TCC CTT TGT TGT GAC TTA TTA AGT CAG GTT Val Ser Leu Arg Ser Phe Ser Leu Cys Cys Asp Leu Leu Ser Gln Val 1490 1495 1500	4692
TGC CAG ACA GCC GTG ACT TAC TGT AAG GAT GCT CTA GAA AAC CAT CTT Cys Gln Thr Ala Val Thr Tyr Cys Lys Asp Ala Leu Glu Asn His Leu 1505 1510 1515	4740
CAT GTT ATT GTT GGT ACA CTT ATA CCC CTT GTG TAT GAG CAG GTG GAG His Val Ile Val Gly Thr Leu Ile Pro Leu Val Tyr Glu Gln Val Glu 1520 1525 1530	4788
GTT CAG AAA CAG GTA TTG GAC TTG TTG AAA TAC TTA GTG ATA GAT AAC Val Gln Lys Gln Val Leu Asp Leu Leu Lys Tyr Leu Val Ile Asp Asn 1535 1540 1545	4836
AAG GAT AAT GAA AAC CTC TAT ATC ACG ATT AAG CTT TTA GAT CCT TTT Lys Asp Asn Glu Asn Leu Tyr Ile Thr Ile Lys Leu Leu Asp Pro Phe 1550 1555 1560 1565	4884
CCT GAC CAT GTT GTT TTT AAG GAT TTG CGT ATT ACT CAG CAA AAA ATC Pro Asp His Val Phe Lys Asp Leu Arg Ile Thr Gln Gln Lys Ile 1570 1575 1580	4932
AAA TAC AGT AGA GGA CCC TTT TCA CTC TTG GAG GAA ATT AAC CAT TTT Lys Tyr Ser Arg Gly Pro Phe Ser Leu Leu Glu Glu Ile Asn His Phe 1585 1590 1595	4980
CTC TCA GTA AGT GTT TAT GAT GCA CTT CCA TTG ACA AGA CTT GAA GGA Leu Ser Val Ser Val Tyr Asp Ala Leu Pro Leu Thr Arg Leu Glu Gly 1600 1605 1610	5028
CTA AAG GAT CTT CGA AGA CAA CTG GAA CTA CAT AAA GAT CAG ATG GTG Leu Lys Asp Leu Arg Arg Gln Leu Glu Leu His Lys Asp Gln Met Val 1615 1620 1625	5076

-188-

GAC ATT ATG AGA GCT TCT CAG GAT AAT CCG CAA GAT GGG ATT ATG GTG Asp Ile Met Arg Ala Ser Gln Asp Asn Pro Gln Asp Gly Ile Met Val 1630 1635 1640 1645	5124
AAA CTA GTT GTC AAT TTG TTG CAG TTA TCC AAG ATG GCA ATA AAC CAC Lys Leu Val Val Asn Leu Leu Gln Leu Ser Lys Met Ala Ile Asn His 1650 1655 1660	5172
ACT GGT GAA AAA GAA GTT CTA GAG GCT GTT GGA AGC TGC TTG GGA GAA Thr Gly Glu Lys Glu Val Leu Glu Ala Val Gly Ser Cys Leu Gly Glu 1665 1670 1675	5220
GTG GGT CCT ATA GAT TTC TCT ACC ATA GCT ATA CAA CAT AGT AAA GAT Val Gly Pro Ile Asp Phe Ser Thr Ile Ala Ile Gln His Ser Lys Asp 1680 1685 1690	5268
GCA TCT TAT ACC AAG GCC CTT AAG TTA TTT GAA GAT AAA GAA CTT CAG Ala Ser Tyr Thr Lys Ala Leu Lys Leu Phe Glu Asp Lys Glu Leu Gln 1695 1700 1705	5316
TGG ACC TTC ATA ATG CTG ACC TAC CTG AAT AAC ACA CTG GTA GAA GAT Trp Thr Phe Ile Met Leu Thr Tyr Leu Asn Asn Thr Leu Val Glu Asp 1710 1715 1720 1725	5364
TGT GTC AAA GTT CGA TCA GCA GCT GTT ACC TGT TTG AAA AAC ATT TTA Cys Val Lys Val Arg Ser Ala Ala Val Thr Cys Leu Lys Asn Ile Leu 1730 1735 1740	5412
GCC ACA AAG ACT GGA CAT AGT TTC TGG GAG ATT TAT AAG ATG ACA ACA Ala Thr Lys Thr Gly His Ser Phe Trp Glu Ile Tyr Lys Met Thr Thr 1745 1750 1755	5460
GAT CCA ATG CTG GCC TAT CTA CAG CCT TTT AGA ACA TCA AGA AAA AAG Asp Pro Met Leu Ala Tyr Leu Gln Pro Phe Arg Thr Ser Arg Lys Lys 1760 1765 1770	5508
TTT TTA GAA GTA CCC AGA TTT GAC AAA GAA AAC CCT TTT GAA GGC CTG Phe Leu Glu Val Pro Arg Phe Asp Lys Glu Asn Pro Phe Glu Gly Leu 1775 1780 1785	5556
GAT GAT ATA AAT CTG TGG ATT CCT CTA AGT GAA AAT CAT GAC ATT TGG Asp Asp Ile Asn Leu Trp Ile Pro Leu Ser Glu Asn His Asp Ile Trp 1790 1795 1800 1805	5604
ATA AAG ACA CTG ACT TGT GCT TTT TTG GAC AGT GGA GGC ACA AAA TGT Ile Lys Thr Leu Thr Cys Ala Phe Leu Asp Ser Gly Gly Thr Lys Cys 1810 1815 1820	5652
GAA ATT CTT CAA TTA TTA AAG CCA ATG TGT GAA GTG AAA ACT GAC TTT Glu Ile Leu Gln Leu Leu Lys Pro Met Cys Glu Val Lys Thr Asp Phe 1825 1830 1835	5700
TGT CAG ACT GTA CTT CCA TAC TTG ATT CAT GAT ATT TTA CTC CAA GAT Cys Gln Thr Val Leu Pro Tyr Leu Ile His Asp Ile Leu Leu Gln Asp 1840 1845 1850	5748
ACA AAT GAA TCA TGG AGA AAT CTG CTT TCT ACA CAT GTT CAG GGA TTT Thr Asn Glu Ser Trp Arg Asn Leu Leu Ser Thr His Val Gln Gly Phe 1855 1860 1865	5796
TTC ACC AGC TGT CTT CGA CAC TTC TCG CAA ACG AGC CGA TCC ACA ACC Phe Thr Ser Cys Leu Arg His Phe Ser Gln Thr Ser Arg Ser Thr Thr 1870 1875 1880 1885	5844
CCT GCA AAC TTG GAT TCA GAG TCA GAG CAC TTT TTC CGA TGC TGT TTG Pro Ala Asn Leu Asp Ser Glu Ser Glu His Phe Phe Arg Cys Cys Leu 1890 1895 1900	5892

-189-

GAT AAA AAA TCA CAA AGA ACA ATG CTT GCT GTT GTG GAC TAC ATG AGA Asp Lys Lys Ser Gln Arg Thr Met Leu Ala Val Val Asp Tyr Met Arg 1905 1910 1915	5940
AGA CAA AAG AGA CCT TCT TCA GGA ACA ATT TTT AAT GAT GCT TTC TGG Arg Gln Lys Arg Pro Ser Ser Gly Thr Ile Phe Asn Asp Ala Phe Trp 1920 1925 1930	5988
CTG GAT TTA AAT TAT CTA GAA GTT GCC AAG GTA GCT CAG TCT TGT GCT Leu Asp Leu Asn Tyr Leu Glu Val Ala Lys Val Ala Gln Ser Cys Ala 1935 1940 1945	6036
GCT CAC TTT ACA GCT TTA CTC TAT GCA GAA ATC TAT GCA GAT AAG AAA Ala His Phe Thr Ala Leu Tyr Ala Glu Ile Tyr Ala Asp Lys Lys 1950 1955 1960 1965	6084
AGT ATG GAT GAT CAA GAG AAA AGA AGT CTT GCA TTT GAA GAA GGA AGC Ser Met Asp Asp Gln Glu Lys Arg Ser Leu Ala Phe Glu Glu Gly Ser 1970 1975 1980	6132
CAG AGT ACA ACT ATT TCT AGC TTG AGT GAA AAA AGT AAA GAA GAA ACT Gln Ser Thr Thr Ile Ser Ser Leu Ser Glu Lys Ser Lys Glu Glu Thr 1985 1990 1995	6180
GGA ATA AGT TTA CAG GAT CTT CTC TTA GAA ATC TAC AGA AGT ATA GGG Gly Ile Ser Leu Gln Asp Leu Leu Leu Glu Ile Tyr Arg Ser Ile Gly 2000 2005 2010	6228
GAG CCA GAT AGT TTG TAT GGC TGT GGT GGA GGG AAG ATG TTA CAA CCC Glu Pro Asp Ser Leu Tyr Gly Cys Gly Gly Gly Lys Met Leu Gln Pro 2015 2020 2025	6276
ATT ACT AGA CTA CGA ACA TAT GAA CAC GAA GCA ATG TGG GGC AAA GCC Ile Thr Arg Leu Arg Thr Tyr Glu His Glu Ala Met Trp Gly Lys Ala 2030 2035 2040 2045	6324
CTA GTA ACA TAT GAC CTC GAA ACA GCA ATC CCC TCA TCA ACA CGC CAG Leu Val Thr Tyr Asp Leu Glu Thr Ala Ile Pro Ser Ser Thr Arg Gln 2050 2055 2060	6372
GCA GGA ATC ATT CAG GCC TTG CAG AAT TTG GGA CTC TGC CAT ATT CTT Ala Gly Ile Ile Gln Ala Leu Gln Asn Leu Gly Leu Cys His Ile Leu 2065 2070 2075	6420
TCC GTC TAT TTA AAA GGA TTG GAT TAT GAA AAT AAA GAC TGG TGT CCT Ser Val Tyr Leu Lys Gly Leu Asp Tyr Glu Asn Lys Asp Trp Cys Pro 2080 2085 2090	6468
GAA CTA GAA GAA CTT CAT TAC CAA GCA GCA TGG AGG AAT ATG CAG TGG Glu Leu Glu Glu Leu His Tyr Gln Ala Ala Trp Arg Asn Met Gln Trp 2095 2100 2105	6516
GAC CAT TGC ACT TCC GTC AGC AAA GAA GTA GAA GGA ACC AGT TAC CAT Asp His Cys Thr Ser Val Ser Lys Glu Val Glu Gly Thr Ser Tyr His 2110 2115 2120 2125	6564
GAA TCA TTG TAC AAT GCT CTA CAA TCT CTA AGA GAC AGA GAA TTC TCT Glu Ser Leu Tyr Asn Ala Leu Gln Ser Leu Arg Asp Arg Glu Phe Ser 2130 2135 2140	6612
ACA TTT TAT GAA AGT CTC AAA TAT GCC AGA GTA AAA GAA GTG GAA GAG Thr Phe Tyr Glu Ser Leu Lys Tyr Ala Arg Val Lys Glu Val Glu Glu 2145 2150 2155	6660
ATG TGT AAG CGC AGC CTT GAG TCT GTG TAT TCG CTC TAT CCC ACA CTT Met Cys Lys Arg Ser Leu Glu Ser Val Tyr Ser Leu Tyr Pro Thr Leu 2160 2165 2170	6708

-190-

AGC	AGG	TTG	CAG	GCC	ATT	GGA	GAG	CTG	GAA	AGC	ATT	GGG	GAG	CTT	TTC	6756
Ser	Arg	Leu	Gln	Ala	Ile	Gly	Glu	Leu	Glu	Ser	Ile	Gly	Glu	Leu	Phe	
	2175					2180					2185					
TCA	AGA	TCA	GTC	ACA	CAT	AGA	CAA	CTC	TCT	GAA	GTA	TAT	ATT	AAG	TGG	6804
Ser	Arg	Ser	Val	Thr	His	Arg	Gln	Leu	Ser	Glu	Val	Tyr	Ile	Lys	Trp	
	2190				2195					2200					2205	
CAG	AAA	CAC	TCC	CAG	CTT	CTC	AAG	GAC	AGT	GAT	TTT	AGT	TTT	CAG	GAG	6852
Gln	Lys	His	Ser	Gln	Leu	Leu	Lys	Asp	Ser	Asp	Phe	Ser	Phe	Gln	Glu	
				2210					2215					2220		
CCT	ATC	ATG	GCT	CTA	CGC	ACA	GTC	ATT	TTG	GAG	ATC	CTG	ATG	GAA	AAG	6900
Pro	Ile	Met	Ala	Leu	Arg	Thr	Val	Ile	Leu	Glu	Ile	Leu	Met	Glu	Lys	
			2225					2230					2235			
GAA	ATG	GAC	AAC	TCA	CAA	AGA	GAA	TGT	ATT	AAG	GAC	ATT	CTC	ACC	AAA	6948
Glu	Met	Asp	Asn	Ser	Gln	Arg	Glu	Cys	Ile	Lys	Asp	Ile	Leu	Thr	Lys	
	2240						2245					2250				
CAC	CTT	GTA	GAA	CTC	TCT	ATA	CTG	GCC	AGA	ACT	TTC	AAG	AAC	ACT	CAG	6996
His	Leu	Val	Glu	Leu	Ser	Ile	Leu	Ala	Arg	Thr	Phe	Lys	Asn	Thr	Gln	
	2255					2260					2265					
CTC	CCT	GAA	AGG	GCA	ATA	TTT	CAA	ATT	AAA	CAG	TAC	AAT	TCA	GTT	AGC	7044
Leu	Pro	Glu	Arg	Ala	Ile	Phe	Gln	Ile	Lys	Gln	Tyr	Asn	Ser	Val	Ser	
	2270				2275					2280					2285	
TGT	GGA	GTC	TCT	GAG	TGG	CAG	CTG	GAA	GAA	GCA	CAA	GTA	TTC	TGG	GCA	7092
Cys	Gly	Val	Ser	Glu	Trp	Gln	Leu	Glu	Glu	Ala	Gln	Val	Phe	Trp	Ala	
				2290				2295						2300		
AAA	AAG	GAG	CAG	AGT	CTT	GCC	CTG	AGT	ATT	CTC	AAG	CAA	ATG	ATC	AAG	7140
Lys	Lys	Glu	Gln	Ser	Leu	Ala	Leu	Ser	Ile	Leu	Lys	Gln	Met	Ile	Lys	
			2305					2310					2315			
AAG	TTG	GAT	GCC	AGC	TGT	GCA	GCG	AAC	AAT	CCC	AGC	CTA	AAA	CTT	ACA	7188
Lys	Leu	Asp	Ala	Ser	Cys	Ala	Ala	Asn	Asn	Pro	Ser	Leu	Lys	Leu	Thr	
	2320					2325						2330				
TAC	ACA	GAA	TGT	CTG	AGG	GTT	TGT	GGC	AAC	TGG	TTA	GCA	GAA	ACG	TGC	7236
Tyr	Thr	Glu	Cys	Leu	Arg	Val	Cys	Gly	Asn	Trp	Leu	Ala	Glu	Thr	Cys	
	2335					2340					2345					
TTA	GAA	AAT	CCT	GCG	GTC	ATC	ATG	CAG	ACC	TAT	CTA	GAA	AAG	GCA	GTA	7284
Leu	Glu	Asn	Pro	Ala	Val	Ile	Met	Gln	Thr	Tyr	Leu	Glu	Lys	Ala	Val	
	2350				2355					2360					2365	
GAA	GTT	GCT	GGA	AAT	TAT	GAT	GGA	GAA	AGT	AGT	GAT	GAG	CTA	AGA	AAT	7332
Glu	Val	Ala	Gly	Asn	Tyr	Asp	Gly	Glu	Ser	Ser	Asp	Glu	Leu	Arg	Asn	
				2370				2375						2380		
GGA	AAA	ATG	AAG	GCA	TTT	CTC	TCA	TTA	GCC	CGG	TTT	TCA	GAT	ACT	CAA	7380
Gly	Lys	Met	Lys	Ala	Phe	Leu	Ser	Leu	Ala	Arg	Phe	Ser	Asp	Thr	Gln	
			2385					2390					2395			
TAC	CAA	AGA	ATT	GAA	AAC	TAC	ATG	AAA	TCA	TCG	GAA	TTT	GAA	AAC	AAG	7428
Tyr	Gln	Arg	Ile	Glu	Asn	Tyr	Met	Lys	Ser	Ser	Glu	Phe	Glu	Asn	Lys	
	2400						2405					2410				
CAA	GCT	CTC	CTG	AAA	AGA	GCC	AAA	GAG	GAA	GTA	GGT	CTC	CTT	AGG	GAA	7476
Gln	Ala	Leu	Leu	Lys	Arg	Ala	Lys	Glu	Glu	Val	Gly	Leu	Leu	Arg	Glu	
	2415					2420					2425					
CAT	AAA	ATT	CAG	ACA	AAC	AGA	TAC	ACA	GTA	AAG	GTT	CAG	CGA	GAG	CTG	7524
His	Lys	Ile	Gln	Thr	Asn	Arg	Tyr	Thr	Val	Lys	Val	Gln	Arg	Glu	Leu	
	2430				2435					2440					2445	

-191-

GAG TTG GAT GAA TTA GCC CTG CGT GCA CTG AAA GAG GAT CGT AAA CGC Glu Leu Asp Glu Leu Ala Leu Arg Ala Leu Lys Glu Asp Arg Lys Arg 2450 2455 2460	7572
TTC TTA TGT AAA GCA GTT GAA AAT TAT ATC AAC TGC TTA TTA AGT GGA Phe Leu Cys Lys Ala Val Glu Asn Tyr Ile Asn Cys Leu Leu Ser Gly 2465 2470 2475	7620
GAA GAA CAT GAT ATG TGG GTA TTC CGG CTT TGT TCC CTC TGG CTT GAA Glu Glu His Asp Met Trp Val Phe Arg Leu Cys Ser Leu Trp Leu Glu 2480 2485 2490	7668
AAT TCT GGA GTT TCT GAA GTC AAT GGC ATG ATG AAG AGA GAC GGA ATG Asn Ser Gly Val Ser Glu Val Asn Gly Met Met Lys Arg Asp Gly Met 2495 2500 2505	7716
AAG ATT CCA ACA TAT AAA TTT TTG CCT CTT ATG TAC CAA TTG GCT GCT Lys Ile Pro Thr Tyr Lys Phe Leu Pro Leu Met Tyr Gln Leu Ala Ala 2510 2515 2520 2525	7764
AGA ATG GGG ACC AAG ATG ATG GGA GGC CTA GGA TTT CAT GAA GTC CTC Arg Met Gly Thr Lys Met Met Gly Gly Leu Gly Phe His Glu Val Leu 2530 2535 2540	7812
AAT AAT CTA ATC TCT AGA ATT TCA ATG GAT CAC CCC CAT CAC ACT TTG Asn Asn Leu Ile Ser Arg Ile Ser Met Asp His Pro His His Thr Leu 2545 2550 2555	7860
TTT ATT ATA CTG GCC TTA GCA AAT GCA AAC AGA GAT GAA TTT CTG ACT Phe Ile Ile Leu Ala Leu Ala Asn Ala Asn Arg Asp Glu Phe Leu Thr 2560 2565 2570	7908
AAA CCA GAG GTA GCC AGA AGA AGC AGA ATA ACT AAA AAT GTG CCT AAA Lys Pro Glu Val Ala Arg Arg Ser Arg Ile Thr Lys Asn Val Pro Lys 2575 2580 2585	7956
CAA AGC TCT CAG CTT GAT GAG GAT CGA ACA GAG GCT GCA AAT AGA ATA Gln Ser Ser Gln Leu Asp Glu Asp Arg Thr Glu Ala Ala Asn Arg Ile 2590 2595 2600 2605	8004
ATA TGT ACT ATC AGA AGT AGG AGA CCT CAG ATG GTC AGA AGT GTT GAG Ile Cys Thr Ile Arg Ser Arg Arg Pro Gln Met Val Arg Ser Val Glu 2610 2615 2620	8052
GCA CTT TGT GAT GCT TAT ATT ATA TTA GCA AAC TTA GAT GCC ACT CAG Ala Leu Cys Asp Ala Tyr Ile Ile Leu Ala Asn Leu Asp Ala Thr Gln 2625 2630 2635	8100
TGG AAG ACT CAG AGA AAA GGC ATA AAT ATT CCA GCA GAC CAG CCA ATT Trp Lys Thr Gln Arg Lys Gly Ile Asn Ile Pro Ala Asp Gln Pro Ile 2640 2645 2650	8148
ACT AAA CTT AAG AAT TTA GAA GAT GTT GTT GTC CCT ACT ATG GAA ATT Thr Lys Leu Lys Asn Leu Glu Asp Val Val Val Pro Thr Met Glu Ile 2655 2660 2665	8196
AAG GTG GAC CAC ACA GGA GAA TAT GGA AAT CTG GTG ACT ATA CAG TCA Lys Val Asp His Thr Gly Glu Tyr Gly Asn Leu Val Thr Ile Gln Ser 2670 2675 2680 2685	8244
TTT AAA GCA GAA TTT CGC TTA GCA GGA GGT GTA AAT TTA CCA AAA ATA Phe Lys Ala Glu Phe Arg Leu Ala Gly Gly Val Asn Leu Pro Lys Ile 2690 2695 2700	8292
ATA GAT TGT GTA GGT TCC GAT GGC AAG GAG AGG AGA CAG CTT GTT AAG Ile Asp Cys Val Gly Ser Asp Gly Lys Glu Arg Arg Gln Leu Val Lys 2705 2710 2715	8340

-192-

GGC	CGT	GAT	GAC	CTG	AGA	CAA	GAT	GCT	GTC	ATG	CAA	CAG	GTC	TTC	CAG	8388
Gly	Arg	Asp	Asp	Leu	Arg	Gln	Asp	Ala	Val	Met	Gln	Gln	Val	Phe	Gln	
		2720					2725					2730				
ATG	TGT	AAT	ACA	TTA	CTG	CAG	AGA	AAC	ACG	GAA	ACT	AGG	AAG	AGG	AAA	8436
Met	Cys	Asn	Thr	Leu	Leu	Gln	Arg	Asn	Thr	Glu	Thr	Arg	Lys	Arg	Lys	
	2735					2740					2745					
TTA	ACT	ATC	TGT	ACT	TAT	AAG	GTG	GTT	CCC	CTC	TCT	CAG	CGA	AGT	GGT	8484
Leu	Thr	Ile	Cys	Thr	Tyr	Lys	Val	Val	Pro	Leu	Ser	Gln	Arg	Ser	Gly	
2750					2755					2760					2765	
GTT	CTT	GAA	TGG	TGC	ACA	GGA	ACT	GTC	CCC	ATT	GGT	GAA	TTT	CTT	GTT	8532
Val	Leu	Glu	Trp	Cys	Thr	Gly	Thr	Val	Pro	Ile	Gly	Glu	Phe	Leu	Val	
				2770					2775					2780		
AAC	AAT	GAA	GAT	GGT	GCT	CAT	AAA	AGA	TAC	AGG	CCA	AAT	GAT	TTC	AGT	8580
Asn	Asn	Glu	Asp	Gly	Ala	His	Lys	Arg	Tyr	Arg	Pro	Asn	Asp	Phe	Ser	
			2785					2790					2795			
GCC	TTT	CAG	TGC	CAA	AAG	AAA	ATG	ATG	GAG	GTG	CAA	AAA	AAG	TCT	TTT	8628
Ala	Phe	Gln	Cys	Gln	Lys	Lys	Met	Met	Glu	Val	Gln	Lys	Lys	Ser	Phe	
		2800					2805					2810				
GAA	GAG	AAA	TAT	GAA	GTC	TTC	ATG	GAT	GTT	TGC	CAA	AAT	TTT	CAA	CCA	8676
Glu	Glu	Lys	Tyr	Glu	Val	Phe	Met	Asp	Val	Cys	Gln	Asn	Phe	Gln	Pro	
	2815					2820					2825					
GTT	TTC	CGT	TAC	TTC	TGC	ATG	GAA	AAA	TTC	TTG	GAT	CCA	GCT	ATT	TGG	8724
Val	Phe	Arg	Tyr	Phe	Cys	Met	Glu	Lys	Phe	Leu	Asp	Pro	Ala	Ile	Trp	
2830					2835					2840					2845	
TTT	GAG	AAG	CGA	TTG	GCT	TAT	ACG	CGC	AGT	GTA	GCT	ACT	TCT	TCT	ATT	8772
Phe	Glu	Lys	Arg	Leu	Ala	Tyr	Thr	Arg	Ser	Val	Ala	Thr	Ser	Ser	Ile	
				2850					2855					2860		
GTT	GGT	TAC	ATA	CTT	GGA	CTT	GGT	GAT	AGA	CAT	GTA	CAG	AAT	ATC	TTG	8820
Val	Gly	Tyr	Ile	Leu	Gly	Leu	Gly	Asp	Arg	His	Val	Gln	Asn	Ile	Leu	
			2865					2870					2875			
ATA	AAT	GAG	CAG	TCA	GCA	GAA	CTT	GTA	CAT	ATA	GAT	CTA	GGT	GTT	GCT	8868
Ile	Asn	Glu	Gln	Ser	Ala	Glu	Leu	Val	His	Ile	Asp	Leu	Gly	Val	Ala	
		2880					2885					2890				
TTT	GAA	CAG	GGC	AAA	ATC	CTT	CCT	ACT	CCT	GAG	ACA	GTT	CCT	TTT	AGA	8916
Phe	Glu	Gln	Gly	Lys	Ile	Leu	Pro	Thr	Pro	Glu	Thr	Val	Pro	Phe	Arg	
	2895					2900					2905					
CTC	ACC	AGA	GAT	ATT	GTG	GAT	GGC	ATG	GGC	ATT	ACG	GGT	GTT	GAA	GGT	8964
Leu	Thr	Arg	Asp	Ile	Val	Asp	Gly	Met	Gly	Ile	Thr	Gly	Val	Glu	Gly	
2910					2915					2920				2925		
GTC	TTC	AGA	AGA	TGC	TGT	GAG	AAA	ACC	ATG	GAA	GTG	ATG	AGA	AAC	TCT	9012
Val	Phe	Arg	Arg	Cys	Cys	Glu	Lys	Thr	Met	Glu	Val	Met	Arg	Asn	Ser	
				2930					2935					2940		
CAG	GAA	ACT	CTG	TTA	ACC	ATT	GTA	GAG	GTC	CTT	CTA	TAT	GAT	CCA	CTC	9060
Gln	Glu	Thr	Leu	Leu	Thr	Ile	Val	Glu	Val	Leu	Leu	Tyr	Asp	Pro	Leu	
			2945					2950					2955			
TTT	GAC	TGG	ACC	ATG	AAT	CCT	TTG	AAA	GCT	TTG	TAT	TTA	CAG	CAG	AGG	9108
Phe	Asp	Trp	Thr	Met	Asn	Pro	Leu	Lys	Ala	Leu	Tyr	Leu	Gln	Gln	Arg	
	2960						2965					2970				
CCG	GAA	GAT	GAA	ACT	GAG	CTT	CAC	CCT	ACT	CTG	AAT	GCA	GAT	GAC	CAA	9156
Pro	Glu	Asp	Glu	Thr	Glu	Leu	His	Pro	Thr	Leu	Asn	Ala	Asp	Asp	Gln	
	2975					2980					2985					

-193-

GAA TGC AAA CGA AAT CTC AGT GAT ATT GAC CAG AGT TTC GAC AAA GTA 9204  
 Glu Cys Lys Arg Asn Leu Ser Asp Ile Asp Gln Ser Phe Asp Lys Val  
 2990 2995 3000 3005

GCT GAA CGT GTC TTA ATG AGA CTA CAA GAG AAA CTG AAA GGA GTG GAA 9252  
 Ala Glu Arg Val Leu Met Arg Leu Gln Glu Lys Leu Lys Gly Val Glu  
 3010 3015 3020

GAA GGC ACT GTG CTC AGT GTT GGT GGA CAG GTG AAT TTG CTC ATA CAG 9300  
 Glu Gly Thr Val Leu Ser Val Gly Gly Gln Val Asn Leu Leu Ile Gln  
 3025 3030 3035

CAG GCC ATA GAC CCC AAA AAT CTC AGC CGA CTT TTC CCA GGA TGG AAA 9348  
 Gln Ala Ile Asp Pro Lys Asn Leu Ser Arg Leu Phe Pro Gly Trp Lys  
 3040 3045 3050

GCT TGG GTG TGATCTTCAG TATATGAATT ACCCTTTC 9385  
 Ala Trp Val  
 3055

## (2) INFORMATION FOR SEQ ID NO:35:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3056 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

Met Ser Leu Val Leu Asn Asp Leu Leu Ile Cys Cys Arg Gln Leu Glu  
 1 5 10 15

His Asp Arg Ala Thr Glu Arg Lys Lys Glu Val Glu Lys Phe Lys Arg  
 20 25 30

Leu Ile Arg Asp Pro Glu Thr Ile Lys His Leu Asp Arg His Ser Asp  
 35 40 45

Ser Lys Gln Gly Lys Tyr Leu Asn Trp Asp Ala Val Phe Arg Phe Leu  
 50 55 60

Gln Lys Tyr Ile Gln Lys Glu Thr Glu Cys Leu Arg Ile Ala Lys Pro  
 65 70 75 80

Asn Val Ser Ala Ser Thr Gln Ala Ser Arg Gln Lys Lys Met Gln Glu  
 85 90 95

Ile Ser Ser Leu Val Lys Tyr Phe Ile Lys Cys Ala Asn Arg Arg Ala  
 100 105 110

Pro Arg Leu Lys Cys Gln Glu Leu Leu Asn Tyr Ile Met Asp Thr Val  
 115 120 125

Lys Asp Ser Ser Asn Gly Ala Ile Tyr Gly Ala Asp Cys Ser Asn Ile  
 130 135 140

Leu Leu Lys Asp Ile Leu Ser Val Arg Lys Tyr Trp Cys Glu Ile Ser  
 145 150 155 160

Gln Gln Gln Trp Leu Glu Leu Phe Ser Val Tyr Phe Arg Leu Tyr Leu  
 165 170 175

Lys Pro Ser Gln Asp Val His Arg Val Leu Val Ala Arg Ile Ile His  
 180 185 190

-194-

Ala	Val	Thr	Lys	Gly	Cys	Cys	Ser	Gln	Thr	Asp	Gly	Leu	Asn	Ser	Lys
	195						200					205			
Phe	Leu	Asp	Phe	Phe	Ser	Lys	Ala	Ile	Gln	Cys	Ala	Arg	Gln	Glu	Lys
	210					215					220				
Ser	Ser	Ser	Gly	Leu	Asn	His	Ile	Leu	Ala	Ala	Leu	Thr	Ile	Phe	Leu
225					230					235					240
Lys	Thr	Leu	Ala	Val	Asn	Phe	Arg	Ile	Arg	Val	Cys	Glu	Leu	Gly	Asp
				245					250					255	
Glu	Ile	Leu	Pro	Thr	Leu	Leu	Tyr	Ile	Trp	Thr	Gln	His	Arg	Leu	Asn
			260					265					270		
Asp	Ser	Leu	Lys	Glu	Val	Ile	Ile	Glu	Leu	Phe	Gln	Leu	Gln	Ile	Tyr
		275					280					285			
Ile	His	His	Pro	Lys	Gly	Ala	Lys	Thr	Gln	Glu	Lys	Gly	Ala	Tyr	Glu
	290					295					300				
Ser	Thr	Lys	Trp	Arg	Ser	Ile	Leu	Tyr	Asn	Leu	Tyr	Asp	Leu	Leu	Val
305					310					315					320
Asn	Glu	Ile	Ser	His	Ile	Gly	Ser	Arg	Gly	Lys	Tyr	Ser	Ser	Gly	Phe
				325					330					335	
Arg	Asn	Ile	Ala	Val	Lys	Glu	Asn	Leu	Ile	Glu	Leu	Met	Ala	Asp	Ile
			340					345					350		
Cys	His	Gln	Val	Phe	Asn	Glu	Asp	Thr	Arg	Ser	Leu	Glu	Ile	Ser	Gln
		355					360					365			
Ser	Tyr	Thr	Thr	Thr	Gln	Arg	Glu	Ser	Ser	Asp	Tyr	Ser	Val	Pro	Cys
		370				375					380				
Lys	Arg	Lys	Lys	Ile	Glu	Leu	Gly	Trp	Glu	Val	Ile	Lys	Asp	His	Leu
385					390					395					400
Gln	Lys	Ser	Gln	Asn	Asp	Phe	Asp	Leu	Val	Pro	Trp	Leu	Gln	Ile	Ala
				405					410					415	
Thr	Gln	Leu	Ile	Ser	Lys	Tyr	Pro	Ala	Ser	Leu	Pro	Asn	Cys	Glu	Leu
			420					425					430		
Ser	Pro	Leu	Leu	Met	Ile	Leu	Ser	Gln	Leu	Leu	Pro	Gln	Gln	Arg	His
		435					440					445			
Gly	Glu	Arg	Thr	Pro	Tyr	Val	Leu	Arg	Cys	Leu	Thr	Glu	Val	Ala	Leu
		450				455						460			
Cys	Gln	Asp	Lys	Arg	Ser	Asn	Leu	Glu	Ser	Ser	Gln	Lys	Ser	Asp	Leu
465					470					475					480
Leu	Lys	Leu	Trp	Asn	Lys	Ile	Trp	Cys	Ile	Thr	Phe	Arg	Gly	Ile	Ser
				485				490						495	
Ser	Glu	Gln	Ile	Gln	Ala	Glu	Asn	Phe	Gly	Leu	Leu	Gly	Ala	Ile	Ile
			500					505					510		
Gln	Gly	Ser	Leu	Val	Glu	Val	Asp	Arg	Glu	Phe	Trp	Lys	Leu	Phe	Thr
		515					520					525			
Gly	Ser	Ala	Cys	Arg	Pro	Ser	Cys	Pro	Ala	Val	Cys	Cys	Leu	Thr	Leu
		530				535					540				



-195-

Ala Leu Thr Thr Ser Ile Val Pro Gly Ala Val Lys Met Gly Ile Glu  
 545 550 555 560  
 Gln Asn Met Cys Glu Val Asn Arg Ser Phe Ser Leu Lys Glu Ser Ile  
 565 570 575  
 Met Lys Trp Leu Leu Phe Tyr Gln Leu Glu Gly Asp Leu Glu Asn Ser  
 580 585 590  
 Thr Glu Val Pro Pro Ile Leu His Ser Asn Phe Pro His Leu Val Leu  
 595 600 605  
 Glu Lys Ile Leu Val Ser Leu Thr Met Lys Asn Cys Lys Ala Ala Met  
 610 615 620  
 Asn Phe Phe Gln Ser Val Pro Glu Cys Glu His His Gln Lys Asp Lys  
 625 630 635 640  
 Glu Glu Leu Ser Phe Ser Glu Val Glu Glu Leu Phe Leu Gln Thr Thr  
 645 650 655  
 Phe Asp Lys Met Asp Phe Leu Thr Ile Val Arg Glu Cys Gly Ile Glu  
 660 665 670  
 Lys His Gln Ser Ser Ile Gly Phe Ser Val His Gln Asn Leu Lys Glu  
 675 680 685  
 Ser Leu Asp Arg Cys Leu Leu Gly Leu Ser Glu Gln Leu Leu Asn Asn  
 690 695 700  
 Tyr Ser Ser Glu Ile Thr Asn Ser Glu Thr Leu Val Arg Cys Ser Arg  
 705 710 715 720  
 Leu Leu Val Gly Val Leu Gly Cys Tyr Cys Tyr Met Gly Val Ile Ala  
 725 730 735  
 Glu Glu Glu Ala Tyr Lys Ser Glu Leu Phe Gln Lys Ala Asn Ser Leu  
 740 745 750  
 Met Gln Cys Ala Gly Glu Ser Ile Thr Leu Phe Lys Asn Lys Thr Asn  
 755 760 765  
 Glu Glu Phe Arg Ile Gly Ser Leu Arg Asn Met Met Gln Leu Cys Thr  
 770 775 780  
 Arg Cys Leu Ser Asn Cys Thr Lys Lys Ser Pro Asn Lys Ile Ala Ser  
 785 790 795 800  
 Gly Phe Phe Leu Arg Leu Leu Thr Ser Lys Leu Met Asn Asp Ile Ala  
 805 810 815  
 Asp Ile Cys Lys Ser Leu Ala Ser Phe Ile Lys Lys Pro Phe Asp Arg  
 820 825 830  
 Gly Glu Val Glu Ser Met Glu Asp Asp Thr Asn Gly Asn Leu Met Glu  
 835 840 845  
 Val Glu Asp Gln Ser Ser Met Asn Leu Phe Asn Asp Tyr Pro Asp Ser  
 850 855 860  
 Ser Val Ser Asp Ala Asn Glu Pro Gly Glu Ser Gln Ser Thr Ile Gly  
 865 870 875 880  
 Ala Ile Asn Pro Leu Ala Glu Glu Tyr Leu Ser Lys Gln Asp Leu Leu  
 885 890 895

-196-

Phe Leu Asp Met Leu Lys Phe Leu Cys Leu Cys Val Thr Thr Ala Gln  
 900 905 910  
 Thr Asn Thr Val Ser Phe Arg Ala Ala Asp Ile Arg Arg Lys Leu Leu  
 915 920 925  
 Met Leu Ile Asp Ser Ser Thr Leu Glu Pro Thr Lys Ser Leu His Leu  
 930 935 940  
 His Met Tyr Leu Met Leu Leu Lys Glu Leu Pro Gly Glu Glu Tyr Pro  
 945 950 955 960  
 Leu Pro Met Glu Asp Val Leu Glu Leu Leu Lys Pro Leu Ser Asn Val  
 965 970 975  
 Cys Ser Leu Tyr Arg Arg Asp Gln Asp Val Cys Lys Thr Ile Leu Asn  
 980 985 990  
 His Val Leu His Val Val Lys Asn Leu Gly Gln Ser Asn Met Asp Ser  
 995 1000 1005  
 Glu Asn Thr Arg Asp Ala Gln Gly Gln Phe Leu Thr Val Ile Gly Ala  
 1010 1015 1020  
 Phe Trp His Leu Thr Lys Glu Arg Lys Tyr Ile Phe Ser Val Arg Met  
 1025 1030 1035 1040  
 Ala Leu Val Asn Cys Leu Lys Thr Leu Leu Glu Ala Asp Pro Tyr Ser  
 1045 1050 1055  
 Lys Trp Ala Ile Leu Asn Val Met Gly Lys Asp Phe Pro Val Asn Glu  
 1060 1065 1070  
 Val Phe Thr Gln Phe Leu Ala Asp Asn His His Gln Val Arg Met Leu  
 1075 1080 1085  
 Ala Ala Glu Ser Ile Asn Arg Leu Phe Gln Asp Thr Lys Gly Asp Ser  
 1090 1095 1100  
 Ser Arg Leu Leu Lys Ala Leu Pro Leu Lys Leu Gln Gln Thr Ala Phe  
 1105 1110 1115 1120  
 Glu Asn Ala Tyr Leu Lys Ala Gln Glu Gly Met Arg Glu Met Ser His  
 1125 1130 1135  
 Ser Ala Glu Asn Pro Glu Thr Leu Asp Glu Ile Tyr Asn Arg Lys Ser  
 1140 1145 1150  
 Val Leu Leu Thr Leu Ile Ala Val Val Leu Ser Cys Ser Pro Ile Cys  
 1155 1160 1165  
 Glu Lys Gln Ala Leu Phe Ala Leu Cys Lys Ser Val Lys Glu Asn Gly  
 1170 1175 1180  
 Leu Glu Pro His Leu Val Lys Lys Val Leu Glu Lys Val Ser Glu Thr  
 1185 1190 1195 1200  
 Phe Gly Tyr Arg Arg Leu Glu Asp Phe Met Ala Ser His Leu Asp Tyr  
 1205 1210 1215  
 Leu Val Leu Glu Trp Leu Asn Leu Gln Asp Thr Glu Tyr Asn Leu Ser  
 1220 1225 1230  
 Ser Phe Pro Phe Ile Leu Leu Asn Tyr Thr Asn Ile Glu Asp Phe Tyr  
 1235 1240 1245

-197-

Arg Ser Cys Tyr Lys Val Leu Ile Pro His Leu Val Ile Arg Ser His  
 1250 1255 1260  
 Phe Asp Glu Val Lys Ser Ile Ala Asn Gln Ile Gln Glu Asp Trp Lys  
 1265 1270 1275 1280  
 Ser Leu Leu Thr Asp Cys Phe Pro Lys Ile Leu Val Asn Ile Leu Pro  
 1285 1290 1295  
 Tyr Phe Ala Tyr Glu Gly Thr Arg Asp Ser Gly Met Ala Gln Gln Arg  
 1300 1305 1310  
 Glu Thr Ala Thr Lys Val Tyr Asp Met Leu Lys Ser Glu Asn Leu Leu  
 1315 1320 1325  
 Gly Lys Gln Ile Asp His Leu Phe Ile Ser Asn Leu Pro Glu Ile Val  
 1330 1335 1340  
 Val Glu Leu Leu Met Thr Leu His Glu Pro Ala Asn Ser Ser Ala Ser  
 1345 1350 1355 1360  
 Gln Ser Thr Asp Leu Cys Asp Phe Ser Gly Asp Leu Asp Pro Ala Pro  
 1365 1370 1375  
 Asn Pro Pro His Phe Pro Ser His Val Ile Lys Ala Thr Phe Ala Tyr  
 1380 1385 1390  
 Ile Ser Asn Cys His Lys Thr Lys Leu Lys Ser Ile Leu Glu Ile Leu  
 1395 1400 1405  
 Ser Lys Ser Pro Asp Ser Tyr Gln Lys Ile Leu Leu Ala Ile Cys Glu  
 1410 1415 1420  
 Gln Ala Ala Glu Thr Asn Asn Val Tyr Lys Lys His Arg Ile Leu Lys  
 1425 1430 1435 1440  
 Ile Tyr His Leu Phe Val Ser Leu Leu Leu Lys Asp Ile Lys Ser Gly  
 1445 1450 1455  
 Leu Gly Gly Ala Trp Ala Phe Val Leu Arg Asp Val Ile Tyr Thr Leu  
 1460 1465 1470  
 Ile His Tyr Ile Asn Gln Arg Pro Ser Cys Ile Met Asp Val Ser Leu  
 1475 1480 1485  
 Arg Ser Phe Ser Leu Cys Cys Asp Leu Leu Ser Gln Val Cys Gln Thr  
 1490 1495 1500  
 Ala Val Thr Tyr Cys Lys Asp Ala Leu Glu Asn His Leu His Val Ile  
 1505 1510 1515 1520  
 Val Gly Thr Leu Ile Pro Leu Val Tyr Glu Gln Val Glu Val Gln Lys  
 1525 1530 1535  
 Gln Val Leu Asp Leu Leu Lys Tyr Leu Val Ile Asp Asn Lys Asp Asn  
 1540 1545 1550  
 Glu Asn Leu Tyr Ile Thr Ile Lys Leu Leu Asp Pro Phe Pro Asp His  
 1555 1560 1565  
 Val Val Phe Lys Asp Leu Arg Ile Thr Gln Gln Lys Ile Lys Tyr Ser  
 1570 1575 1580  
 Arg Gly Pro Phe Ser Leu Leu Glu Glu Ile Asn His Phe Leu Ser Val  
 1585 1590 1595 1600

-198-

Ser Val Tyr Asp Ala Leu Pro Leu Thr Arg Leu Glu Gly Leu Lys Asp  
 1605 1610 1615  
 Leu Arg Arg Gln Leu Glu Leu His Lys Asp Gln Met Val Asp Ile Met  
 1620 1625 1630  
 Arg Ala Ser Gln Asp Asn Pro Gln Asp Gly Ile Met Val Lys Leu Val  
 1635 1640 1645  
 Val Asn Leu Leu Gln Leu Ser Lys Met Ala Ile Asn His Thr Gly Glu  
 1650 1655 1660  
 Lys Glu Val Leu Glu Ala Val Gly Ser Cys Leu Gly Glu Val Gly Pro  
 1665 1670 1675 1680  
 Ile Asp Phe Ser Thr Ile Ala Ile Gln His Ser Lys Asp Ala Ser Tyr  
 1685 1690 1695  
 Thr Lys Ala Leu Lys Leu Phe Glu Asp Lys Glu Leu Gln Trp Thr Phe  
 1700 1705 1710  
 Ile Met Leu Thr Tyr Leu Asn Asn Thr Leu Val Glu Asp Cys Val Lys  
 1715 1720 1725  
 Val Arg Ser Ala Ala Val Thr Cys Leu Lys Asn Ile Leu Ala Thr Lys  
 1730 1735 1740  
 Thr Gly His Ser Phe Trp Glu Ile Tyr Lys Met Thr Thr Asp Pro Met  
 1745 1750 1755 1760  
 Leu Ala Tyr Leu Gln Pro Phe Arg Thr Ser Arg Lys Lys Phe Leu Glu  
 1765 1770 1775  
 Val Pro Arg Phe Asp Lys Glu Asn Pro Phe Glu Gly Leu Asp Asp Ile  
 1780 1785 1790  
 Asn Leu Trp Ile Pro Leu Ser Glu Asn His Asp Ile Trp Ile Lys Thr  
 1795 1800 1805  
 Leu Thr Cys Ala Phe Leu Asp Ser Gly Gly Thr Lys Cys Glu Ile Leu  
 1810 1815 1820  
 Gln Leu Leu Lys Pro Met Cys Glu Val Lys Thr Asp Phe Cys Gln Thr  
 1825 1830 1835 1840  
 Val Leu Pro Tyr Leu Ile His Asp Ile Leu Leu Gln Asp Thr Asn Glu  
 1845 1850 1855  
 Ser Trp Arg Asn Leu Leu Ser Thr His Val Gln Gly Phe Phe Thr Ser  
 1860 1865 1870  
 Cys Leu Arg His Phe Ser Gln Thr Ser Arg Ser Thr Thr Pro Ala Asn  
 1875 1880 1885  
 Leu Asp Ser Glu Ser Glu His Phe Phe Arg Cys Cys Leu Asp Lys Lys  
 1890 1895 1900  
 Ser Gln Arg Thr Met Leu Ala Val Val Asp Tyr Met Arg Arg Gln Lys  
 1905 1910 1915 1920  
 Arg Pro Ser Ser Gly Thr Ile Phe Asn Asp Ala Phe Trp Leu Asp Leu  
 1925 1930 1935  
 Asn Tyr Leu Glu Val Ala Lys Val Ala Gln Ser Cys Ala Ala His Phe  
 1940 1945 1950

-199-

Thr Ala Leu Leu Tyr Ala Glu Ile Tyr Ala Asp Lys Lys Ser Met Asp  
 1955 1960 1965  
 Asp Gln Glu Lys Arg Ser Leu Ala Phe Glu Glu Gly Ser Gln Ser Thr  
 1970 1975 1980  
 Thr Ile Ser Ser Leu Ser Glu Lys Ser Lys Glu Glu Thr Gly Ile Ser  
 1985 1990 1995 2000  
 Leu Gln Asp Leu Leu Leu Glu Ile Tyr Arg Ser Ile Gly Glu Pro Asp  
 2005 2010 2015  
 Ser Leu Tyr Gly Cys Gly Gly Gly Lys Met Leu Gln Pro Ile Thr Arg  
 2020 2025 2030  
 Leu Arg Thr Tyr Glu His Glu Ala Met Trp Gly Lys Ala Leu Val Thr  
 2035 2040 2045  
 Tyr Asp Leu Glu Thr Ala Ile Pro Ser Ser Thr Arg Gln Ala Gly Ile  
 2050 2055 2060  
 Ile Gln Ala Leu Gln Asn Leu Gly Leu Cys His Ile Leu Ser Val Tyr  
 2065 2070 2075 2080  
 Leu Lys Gly Leu Asp Tyr Glu Asn Lys Asp Trp Cys Pro Glu Leu Glu  
 2085 2090 2095  
 Glu Leu His Tyr Gln Ala Ala Trp Arg Asn Met Gln Trp Asp His Cys  
 2100 2105 2110  
 Thr Ser Val Ser Lys Glu Val Glu Gly Thr Ser Tyr His Glu Ser Leu  
 2115 2120 2125  
 Tyr Asn Ala Leu Gln Ser Leu Arg Asp Arg Glu Phe Ser Thr Phe Tyr  
 2130 2135 2140  
 Glu Ser Leu Lys Tyr Ala Arg Val Lys Glu Val Glu Glu Met Cys Lys  
 2145 2150 2155 2160  
 Arg Ser Leu Glu Ser Val Tyr Ser Leu Tyr Pro Thr Leu Ser Arg Leu  
 2165 2170 2175  
 Gln Ala Ile Gly Glu Leu Glu Ser Ile Gly Glu Leu Phe Ser Arg Ser  
 2180 2185 2190  
 Val Thr His Arg Gln Leu Ser Glu Val Tyr Ile Lys Trp Gln Lys His  
 2195 2200 2205  
 Ser Gln Leu Leu Lys Asp Ser Asp Phe Ser Phe Gln Glu Pro Ile Met  
 2210 2215 2220  
 Ala Leu Arg Thr Val Ile Leu Glu Ile Leu Met Glu Lys Glu Met Asp  
 2225 2230 2235 2240  
 Asn Ser Gln Arg Glu Cys Ile Lys Asp Ile Leu Thr Lys His Leu Val  
 2245 2250 2255  
 Glu Leu Ser Ile Leu Ala Arg Thr Phe Lys Asn Thr Gln Leu Pro Glu  
 2260 2265 2270  
 Arg Ala Ile Phe Gln Ile Lys Gln Tyr Asn Ser Val Ser Cys Gly Val  
 2275 2280 2285  
 Ser Glu Trp Gln Leu Glu Glu Ala Gln Val Phe Trp Ala Lys Lys Glu  
 2290 2295 2300

-200-

Gln Ser Leu Ala Leu Ser Ile Leu Lys Gln Met Ile Lys Lys Leu Asp  
 2305 2310 2315 2320  
 Ala Ser Cys Ala Ala Asn Asn Pro Ser Leu Lys Leu Thr Tyr Thr Glu  
 2325 2330 2335  
 Cys Leu Arg Val Cys Gly Asn Trp Leu Ala Glu Thr Cys Leu Glu Asn  
 2340 2345 2350  
 Pro Ala Val Ile Met Gln Thr Tyr Leu Glu Lys Ala Val Glu Val Ala  
 2355 2360 2365  
 Gly Asn Tyr Asp Gly Glu Ser Ser Asp Glu Leu Arg Asn Gly Lys Met  
 2370 2375 2380  
 Lys Ala Phe Leu Ser Leu Ala Arg Phe Ser Asp Thr Gln Tyr Gln Arg  
 2385 2390 2395 2400  
 Ile Glu Asn Tyr Met Lys Ser Ser Glu Phe Glu Asn Lys Gln Ala Leu  
 2405 2410 2415  
 Leu Lys Arg Ala Lys Glu Glu Val Gly Leu Leu Arg Glu His Lys Ile  
 2420 2425 2430  
 Gln Thr Asn Arg Tyr Thr Val Lys Val Gln Arg Glu Leu Glu Leu Asp  
 2435 2440 2445  
 Glu Leu Ala Leu Arg Ala Leu Lys Glu Asp Arg Lys Arg Phe Leu Cys  
 2450 2455 2460  
 Lys Ala Val Glu Asn Tyr Ile Asn Cys Leu Leu Ser Gly Glu Glu His  
 2465 2470 2475 2480  
 Asp Met Trp Val Phe Arg Leu Cys Ser Leu Trp Leu Glu Asn Ser Gly  
 2485 2490 2495  
 Val Ser Glu Val Asn Gly Met Met Lys Arg Asp Gly Met Lys Ile Pro  
 2500 2505 2510  
 Thr Tyr Lys Phe Leu Pro Leu Met Tyr Gln Leu Ala Ala Arg Met Gly  
 2515 2520 2525  
 Thr Lys Met Met Gly Gly Leu Gly Phe His Glu Val Leu Asn Asn Leu  
 2530 2535 2540  
 Ile Ser Arg Ile Ser Met Asp His Pro His His Thr Leu Phe Ile Ile  
 2545 2550 2555 2560  
 Leu Ala Leu Ala Asn Ala Asn Arg Asp Glu Phe Leu Thr Lys Pro Glu  
 2565 2570 2575  
 Val Ala Arg Arg Ser Arg Ile Thr Lys Asn Val Pro Lys Gln Ser Ser  
 2580 2585 2590  
 Gln Leu Asp Glu Asp Arg Thr Glu Ala Ala Asn Arg Ile Ile Cys Thr  
 2595 2600 2605  
 Ile Arg Ser Arg Arg Pro Gln Met Val Arg Ser Val Glu Ala Leu Cys  
 2610 2615 2620  
 Asp Ala Tyr Ile Ile Leu Ala Asn Leu Asp Ala Thr Gln Trp Lys Thr  
 2625 2630 2635 2640  
 Gln Arg Lys Gly Ile Asn Ile Pro Ala Asp Gln Pro Ile Thr Lys Leu  
 2645 2650 2655

-201-

Lys Asn Leu Glu Asp Val Val Val Pro Thr Met Glu Ile Lys Val Asp  
 2660 2665 2670  
 His Thr Gly Glu Tyr Gly Asn Leu Val Thr Ile Gln Ser Phe Lys Ala  
 2675 2680 2685  
 Glu Phe Arg Leu Ala Gly Gly Val Asn Leu Pro Lys Ile Ile Asp Cys  
 2690 2695 2700  
 Val Gly Ser Asp Gly Lys Glu Arg Arg Gln Leu Val Lys Gly Arg Asp  
 2705 2710 2715 2720  
 Asp Leu Arg Gln Asp Ala Val Met Gln Gln Val Phe Gln Met Cys Asn  
 2725 2730 2735  
 Thr Leu Leu Gln Arg Asn Thr Glu Thr Arg Lys Arg Lys Leu Thr Ile  
 2740 2745 2750  
 Cys Thr Tyr Lys Val Val Pro Leu Ser Gln Arg Ser Gly Val Leu Glu  
 2755 2760 2765  
 Trp Cys Thr Gly Thr Val Pro Ile Gly Glu Phe Leu Val Asn Asn Glu  
 2770 2775 2780  
 Asp Gly Ala His Lys Arg Tyr Arg Pro Asn Asp Phe Ser Ala Phe Gln  
 2785 2790 2795 2800  
 Cys Gln Lys Lys Met Met Glu Val Gln Lys Lys Ser Phe Glu Glu Lys  
 2805 2810 2815  
 Tyr Glu Val Phe Met Asp Val Cys Gln Asn Phe Gln Pro Val Phe Arg  
 2820 2825 2830  
 Tyr Phe Cys Met Glu Lys Phe Leu Asp Pro Ala Ile Trp Phe Glu Lys  
 2835 2840 2845  
 Arg Leu Ala Tyr Thr Arg Ser Val Ala Thr Ser Ser Ile Val Gly Tyr  
 2850 2855 2860  
 Ile Leu Gly Leu Gly Asp Arg His Val Gln Asn Ile Leu Ile Asn Glu  
 2865 2870 2875 2880  
 Gln Ser Ala Glu Leu Val His Ile Asp Leu Gly Val Ala Phe Glu Gln  
 2885 2890 2895  
 Gly Lys Ile Leu Pro Thr Pro Glu Thr Val Pro Phe Arg Leu Thr Arg  
 2900 2905 2910  
 Asp Ile Val Asp Gly Met Gly Ile Thr Gly Val Glu Gly Val Phe Arg  
 2915 2920 2925  
 Arg Cys Cys Glu Lys Thr Met Glu Val Met Arg Asn Ser Gln Glu Thr  
 2930 2935 2940  
 Leu Leu Thr Ile Val Glu Val Leu Leu Tyr Asp Pro Leu Phe Asp Trp  
 2945 2950 2955 2960  
 Thr Met Asn Pro Leu Lys Ala Leu Tyr Leu Gln Gln Arg Pro Glu Asp  
 2965 2970 2975  
 Glu Thr Glu Leu His Pro Thr Leu Asn Ala Asp Asp Gln Glu Cys Lys  
 2980 2985 2990  
 Arg Asn Leu Ser Asp Ile Asp Gln Ser Phe Asp Lys Val Ala Glu Arg  
 2995 3000 3005

-202-

Val Leu Met Arg Leu Gln Glu Lys Leu Lys Gly Val Glu Glu Gly Thr  
 3010 3015 3020

Val Leu Ser Val Gly Gly Gln Val Asn Leu Leu Ile Gln Gln Ala Ile  
 3025 3030 3035 3040

Asp Pro Lys Asn Leu Ser Arg Leu Phe Pro Gly Trp Lys Ala Trp Val  
 3045 3050 3055

## (2) INFORMATION FOR SEQ ID NO:36:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 19 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

Met Ser Gly Gly Ser Ser Cys Gln Thr Pro Ser Arg Ala Ile Pro Ala  
 1 5 10 15  
 Thr Arg Arg

## (2) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 21 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

Gly Asp Tyr Ser Thr Thr Pro Gly Gly Thr Leu Phe Ser Thr Thr Pro  
 1 5 10 15  
 Gly Gly Thr Arg Arg  
 20

## (2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 12 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

Glu Cys Arg Asn Ser Pro Val Thr Lys Thr Arg Arg  
 1 5 10



-203-

## (2) INFORMATION FOR SEQ ID NO:39:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 12 amino acids  
    (B) TYPE: amino acid  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

Gly Val Thr Ser Pro Ser Ser Asp Glu Pro Arg Arg  
1                    5                    10

## (2) INFORMATION FOR SEQ ID NO:40:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 10 amino acids  
    (B) TYPE: amino acid  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

Met Glu Ala Ser Gln Ser His Leu Arg Arg  
1                    5                    10

## (2) INFORMATION FOR SEQ ID NO:41:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 12 amino acids  
    (B) TYPE: amino acid  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

Arg Arg Asn Ser Pro Glu Asp Lys Arg Ala Gly Gly  
1                    5                    10

## (2) INFORMATION FOR SEQ ID NO:42:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 12 amino acids  
    (B) TYPE: amino acid  
    (D) TOPOLOGY: linear

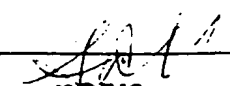
(ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

Gly Glu Glu Ser Gln Phe Glu Met Asp Ile Arg Arg  
1                    5                    10

## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>10</u> , lines <u>2-6</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, MD 20852 US	
Date of deposit 7 November 1996	Accession Number S HB 12233 and HB 12234
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>"In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC)."</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
EP	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	
For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer  International Division (703) 295-3880	Authorized officer

-205-

**CLAIMS**

We claim:

1. A purified and isolated polynucleotide comprising a polynucleotide encoding the PIK-related kinase MCCS1 $\alpha$  amino acid sequence set out in SEQ ID NO: 31.
2. A purified and isolated polynucleotide comprising a polynucleotide encoding the PIK-related kinase MCCS1 $\beta$  amino acid sequence set out in SEQ ID NO: 33.
3. The polynucleotide of claim 1 or 2 which is a DNA.
4. The DNA of claim 3 which is a cDNA.
5. A MCCS1 $\alpha$  cDNA consisting of the DNA sequence set out in SEQ ID NO: 30.
6. A MCCS1 $\beta$  DNA consisting of the DNA sequence set out in SEQ ID NO: 32.
7. The DNA of claim 3 which is a genomic DNA.
8. An RNA transcript of the DNA of claim 3.
9. The DNA of claim 3 which is a wholly or partially chemically synthesized DNA.

-206-

10. A DNA comprising a DNA encoding a full length mammalian MCCS1 kinase selected from the group consisting of:

- a) a DNA which hybridizes under stringent conditions to the non-coding strand of the DNA of SEQ ID NO: 30;
- b) a DNA which hybridizes under stringent conditions to the non-coding strand of the DNA of SEQ ID NO: 3; and
- c) a DNA which hybridizes under stringent conditions to the non-coding strand of the DNA of SEQ ID NO: 32.

11. A vector comprising a DNA according to claim 3 or 10.

12. The vector of claim 11 wherein said DNA is operatively linked to an expression control DNA sequence.

13. A host cell stably transformed or transfected with a DNA according to claim 3 or 10 in a manner allowing the expression in said host cell of the MCCS1 kinase.

14. A method for producing the PIK-related kinase MCCS1, said method comprising growing a host cell according to claim 11 in a suitable nutrient medium and isolating the MCCS1 kinase from said cell or the medium of its growth.

15. A purified and isolated polypeptide comprising the PIK-related kinase MCCS1 $\alpha$  amino acid sequence consisting of SEQ ID NO: 31.

16. A purified and isolated polypeptide comprising the PIK-related kinase MCCS1 $\beta$  amino acid sequence consisting of SEQ ID NO: 33.

17. A polypeptide or peptide capable of specifically binding to PIK-related kinase MCCS1.

18. An antibody product according to claim 17.

-207-

19. A monoclonal antibody according to claim 18.
20. A hybridoma cell line producing a monoclonal antibody according to claim 19.
21. An assay for identifying modulators of MCCS1 kinase activity comprising the steps of:
  - a) incubating a MCCS1 kinase preparation in kinase buffer with gamma-<sup>32</sup>P-ATP and an exogenous kinase substrate in the presence and absence of a test compound, and
  - b) measuring the moles of phosphate transferred to said substrate; wherein an increase in the moles of <sup>32</sup>P-phosphate transferred to said substrate in presence of said test compound compared to the moles of <sup>32</sup>P-phosphate transferred to said substrate in the absence of said test compound indicates that said test compound is an activator of said MCCS1 kinase and a decrease in the moles of <sup>32</sup>P-phosphate transferred to said substrate in presence of said test compound compared to the moles of <sup>32</sup>P-phosphate transferred to said substrate in the absence of said test compound indicates that said test compound is an inhibitor of said MCCS1 kinase.
22. The hybridoma cell line 224C.
23. The hybridoma cell line 224F.

-208-

24. A method of identifying a compound that inhibits MCCC1 comprising the steps of:

- a) expressing MCCC1 in a genetically altered cell, thereby decreasing the sensitivity of the cell to DNA damage, said sensitivity being associated with the genetic alteration;
- b) exposing the genetically altered cell of step (a) to DNA damaging treatment in the presence and absence of a test modulator compound;
- c) measuring the sensitivity of the cell to DNA damage; and
- d) identifying a test compound that restores the sensitivity of the cell to DNA damage as an inhibitor of MCCC1 activity.

25. A method of identifying a compound that inhibits ATM comprising the steps of:

- a) expressing ATM in a genetically altered cell, thereby decreasing the sensitivity of the cell to DNA damage, said sensitivity being associated with the genetic alteration;
- b) exposing the genetically altered cell of step (a) to DNA damaging treatment in the presence and absence of a test modulator compound;
- c) measuring the sensitivity of the cell to DNA damage; and
- d) identifying a test compound that restores the sensitivity of the cell to DNA damage as an inhibitor of ATM activity.

-209-

26. An assay for identifying modulators of ATM kinase activity comprising the steps of:

a) incubating a ATM kinase preparation in kinase buffer with gamma-<sup>32</sup>P-ATP and an exogenous kinase substrate in the presence and absence of a test compound, and

b) measuring the moles of phosphate transferred to said substrate; wherein an increase in the moles of <sup>32</sup>P-phosphate transferred to said substrate in presence of said test compound compared to the moles of <sup>32</sup>P-phosphate transferred to said substrate in the absence of said test compound indicates that said test compound is an activator of said ATM kinase and a decrease in the moles of <sup>32</sup>P-phosphate transferred to said substrate in presence of said test compound compared to the moles of <sup>32</sup>P-phosphate transferred to said substrate in the absence of said test compound indicates that said test compound is an inhibitor of said ATM kinase.





<b>(51) International Patent Classification <sup>6</sup> :</b> <b>C12N 9/12, 15/63, C07K 16/40, C12N</b> <b>5/12, C12O 1/48, C12N 15/11</b>	<b>A3</b>	<b>(11) International Publication Number: WO 97/18323</b>
		<b>(43) International Publication Date: 22 May 1997 (22.05.97)</b>

The present invention generally relates to genes encoding cell cycle checkpoint phosphatidylinositol kinase (PIK)-related proteins essential to DNA damage responses in cells. These PIK-related kinases are required in regulatory pathways that arrest the cell cycle following DNA damage to allow DNA repair prior to mitosis or initiation of DNA replication. More particularly, the invention provides a novel human cell cycle checkpoint PIK-related kinase, MCCC1, and polynucleotide sequences encoding the MCCC1. Assays for identifying modulators of MCCC1 useful as, for example, chemotherapy and radiation adjuvants, are also provided by the invention. Further, assays for identifying modulators of the cell cycle checkpoint phosphatidylinositol kinase (PIK)-related protein identified as ATM are provided.

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 96/19337

<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC 6 C12N9/12 C12N15/63 C07K16/40 C12N5/12 C12Q1/48 C12N15/11		
According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) IPC 6 C12N		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 97 09433 A (MEDICAL RESEARCH COUNCIL) 13 March 1997  see SEQ ID No. 1 and 2; pages 16-22; page 33, lines 28-33; page 34, lines 1-12 ---	2-4, 9-14, 16-18, 21,24
P,X	PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES, vol. 93, April 1996, pages 2850-2855, XP002023632 CIMPRICH, K.A. ET AL.: "cDNA cloning and gene mapping of a candidate human cell cycle checkpoint protein" see Figure 2 --- <div style="text-align: center;">-/-</div>	1,3,4, 9-11,15, 16
<div style="display: flex; justify-content: space-between;"> <span><input checked="" type="checkbox"/> Further documents are listed in the continuation of box C.</span> <span><input checked="" type="checkbox"/> Patent family members are listed in annex.</span> </div>		
* Special categories of cited documents : <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>*A* document defining the general state of the art which is not considered to be of particular relevance</p> <p>*E* earlier document but published on or after the international filing date</p> <p>*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>*O* document referring to an oral disclosure, use, exhibition or other means</p> <p>*P* document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>*Z* document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search	Date of mailing of the international search report	
1 August 1997	26.08.97	
Name and mailing address of the ISA	Authorized officer	
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl, Fax (+ 31-70) 340-3016	Alt, G	

# INTERNATIONAL SEARCH REPORT

National Application No.

PCT/US 96/19337

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>GENE, vol. 119, 1992, pages 83-89, XP002023633 SEATON, B.L. ET AL.: "Isolation and characterization of the Schizosaccharomyces pombe rad3 gene, involved in the DNA damage and DNA synthesis checkpoints" cited in the application see Figure 5; page 84, right-hand column ---</p>	1
P,A	<p>EMBO JOURNAL, vol. 15, no. 23, 1996, pages 6641-6651, XP002023634 BENTLEY, N.J. ET AL.: "The Schizosaccharomyces pombe rad3 checkpoint gene" see the whole document ---</p>	1
A	<p>SCIENCE, vol. 268, 23 June 1995, pages 1749-1753, XP002036686 SAVITSKY, K. ET AL.: "A single ataxia telangiectasia gene with a product similar to PI-3 kinase" see the whole document ---</p>	25,26
T	<p>CURRENT OPINION GENET. DEV., vol. 7, no. 2, 1997, pages 170-175, XP002036687 HOEKSTRA, M.F.: "Responses to DNA damage and regulation of cell cycle checkpoints by the ATM protein kinase family" see the whole document -----</p>	1



